

# G.I. FORUM

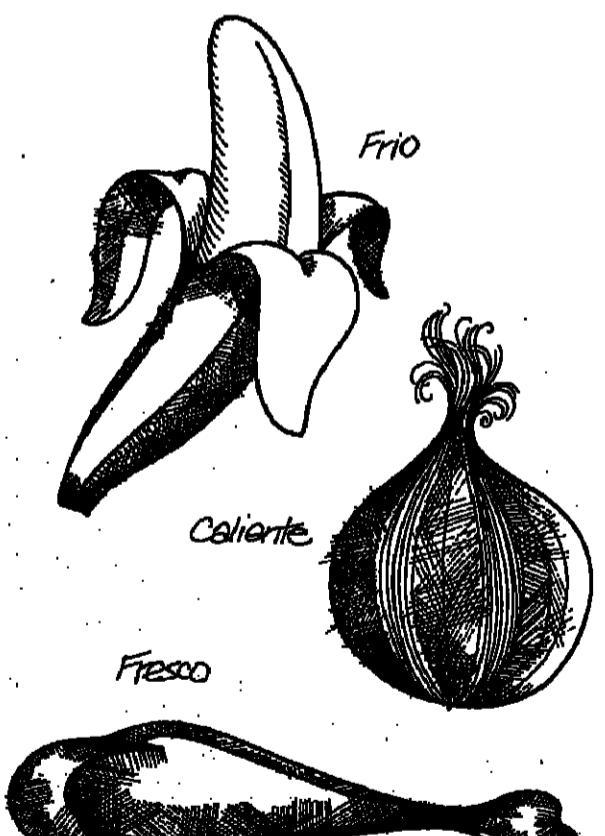
A CURRENT REVIEW OF INVESTIGATIONS IN GASTROENTEROLOGY



## Duodenal ulcer treatment and the "hot-cold" theory



A recent ethnographic study of a group of Spanish-speaking residents in New York City revealed an ancient "hot-cold" theory of disease not only still prevalent, but also compatible with some aspects of current ulcer management.<sup>1</sup>



The theory stems from the classic Greek humoral system of disease which was transferred to the New World by the Spanish and Portuguese in the 16th and 17th centuries. In the variant studied in New York, diseases and bodily conditions are classified as either "hot" (*caliente*) or "cold" (*frio*) and foods and medicines as "hot," "cold" or "cool."

(*fresco*), irrespective of their actual temperatures. According to the theory, a "hot" condition should be treated with "cold" foods and medicines, and vice versa.<sup>1</sup>

Sometimes this presents a problem in modern medical management. For example, pregnant women often refuse "hot" iron supplements or vitamins in order to prevent their babies from being born with a rash, a "hot" condition. But in ulcer—another "hot" condition—the bland diet, still so frequently prescribed today, prohibits most of the foods considered "hot" within the folk system, including spices and coffee.<sup>1</sup>

## Milk, chicken breast—and horseradish?

However, the bland diet itself now tends to be considered in many quarters almost akin to folk medicine. One investigator notes that since the time of the 19th century French pathologist Jean Cruveilhier, the bland diet has been synonymous with the "white" diet—milk, chicken breast, cottage cheese. But what about white horseradish? he wonders. His point—much of dietotherapy by analogy may be ludicrous.<sup>2</sup>

## Milk, antacid and hospitalization

Further thrust to this argument was given by controlled studies alternating an unrestricted diet with a standard bland diet in patients diagnosed as having active duodenal ulcer. One such study, in Iowa, showed no significant difference in healing rates, symptoms or recurrences between patients given a bland diet and those given a standard one.<sup>3</sup>

A British observer<sup>4</sup> states that while these results suggest diet has no effect on the remission of duodenal ulcer, they do not constitute absolute proof. To begin with, all of the patients were given regular and frequent doses of milk and antacids. But most important of all, they were hospitalized for purposes of the study. And hospitalization alone is known to bring relief to the ulcer patient.<sup>4</sup>

References: 1. Harwood, A.: *J.A.M.A.*, 216:1152, 1971; 2. Ingelfinger, F. J.: "Let the Ulcer Patient Enjoy His Food," in Ingelfinger, F. J.; Relman, A. S., and Finland, M. (eds.): *Controversy in Internal Medicine*, Philadelphia, W. B. Saunders Co., 1966, p. 173. 3. Buchman, E., et al.: *Gastroenterology*, 56:1016, 1969. 4. Diet and Duodenal Ulcer, *Brit. Med. J.*, 3:727, 1969.

## Librax®—for excessive anxiety and related G.I. symptoms

Excessive anxiety can be a major triggering stimulus, inducing gastrointestinal hypersecretion and hypermotility and frequently leading to ulcer exacerbation in a susceptible individual. For many duodenal ulcer patients hospitalization may be unwarranted, long vacations impractical—but they still need respite from hypermotility and hypersecretion which produce spasm and associated pain. In many cases, adjunctive Librax can help. Only Librax offers in a single capsule the well-known antianxiety action of Librium® (chlordiazepoxide HCl) and the antisecretory/antispasmodic action of Quarzan® (clidinium Br).

## The logic of dual-action therapy

The action of Librium usually helps reduce excessive anxiety which may accentuate the somatic symptoms of duodenal ulcer. At the same time, the action of Quarzan, a dependable anticholinergic, helps reduce gastric hypersecretion and hypermotility—thereby helping to relieve spasm and associated pain.

While the evidence is inconclusive regarding the precise role dietotherapy may play in gastroenterologic medicine, the value of adjunctive Librax in the total medical management of the peptic duodenal ulcer patient has been clearly demonstrated.

## Up to 8 capsules daily in divided doses

For optimum response, dosage may be adjusted to your patients' requirements, within the range of 1 or 2 capsules, 3 or 4 times daily.

Before prescribing, please consult complete product information, a summary of which follows:

**Indications:** Symptomatic relief of hypersecretion, hypermotility and anxiety and tension states associated with organic or functional gastrointestinal disorders; and as adjunctive therapy in the management of peptic ulcer, gastritis, duodenitis, irritable bowel syndrome, spastic colitis, and mild ulcerative colitis.

**Contraindications:** Patients with glaucoma; prostate hypertrophy and benign bladder neck obstruction; known hypersensitivity to chlordiazepoxide hydrochloride and/or clidinium bromide.

**Warnings:** Caution patients about possible combined effects with alcohol and other CNS depressants. As with all CNS-acting drugs, caution patients against hazardous occupations requiring complete mental alertness (e.g., operating machinery, driving). Though physical and psychological dependence have rarely been reported on recommended doses, use caution in administering Librium (chlordiazepoxide hydrochloride) to known addiction-prone individuals or those who might increase dosage; withdrawn symptoms (including convulsions), following discontinuation of the drug and similar to those seen with barbiturates, have been reported. Use of any drug in pregnancy, lactation, or in women of childbearing age requires that its potential benefits be weighed against its possible hazards. As with all anticholinergic drugs, an inhibiting effect on lactation may occur.

**Precautions:** In elderly and debilitated, limit dosage to smallest effective amount to preclude development of ataxia, oversedation or confusion (not more than two capsules per day initially; increase gradually as needed and tolerated). Though generally not recommended, if combination therapy with other psychotropics seems indicated, carefully consider individual pharmacologic effects, particularly in use of potentiating drugs such as MAO inhibitors and phenothiazines. Observe usual precautions in presence of impaired renal or hepatic function. Paradoxical reactions (e.g., excitement, stimulation and acute rage) have been reported in psychiatric patients. Employ usual precautions in treatment of anxiety states with evidence of impending depression; suicidal tendencies may be present and protective measures necessary. Variable effects on blood coagulation have been reported very rarely in patients receiving the drug and antiocoagulants; causal relationship has not been established clinically.

**Adverse Reactions:** No side effects or manifestations not seen with either compound alone have been reported with Librax. When chlordiazepoxide hydrochloride is used alone, drowsiness, ataxia and confusion may occur, especially in the elderly and debilitated. These are reversible in most instances by proper dosage adjustment, but are also occasionally observed at the lower dosage ranges. In a few instances syncope has been reported. Also encountered are isolated instances of skin eruptions, edema, minor menstrual irregularities, nausea and constipation, extrapyramidal symptoms, increased and decreased libido—all infrequent and generally controlled with dosage reduction; changes in EEG patterns (low-voltage fast activity) may appear during and after therapy; blood dyscrasias (including agranulocytosis), jaundice and hepatic dysfunction have been reported, occasionally with chlordiazepoxide hydrochloride, making periodic blood counts and liver function tests advisable during protracted therapy. Adverse effects reported with Librax are typical of anticholinergic agents, i.e., dryness of mouth, blurring of vision, urinary hesitancy and constipation. Constipation has occurred most often when Librax therapy is combined with other spasmolytics and/or low residue diets.

They added that the relabeling may prove

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## Jackson Is Stonewalled

## Amphetamine Regimen Calms Vicious Dog

Medical Tribune Report

WASHINGTON—The case history of a hyperkinetic dog whose extreme violence and viciousness disappeared within an hour after dextroamphetamine therapy and has not recurred was outlined here by an Ohio investigator during the annual meeting of the American Association for the Advancement of Science.

The experiment suggests that certain drugs may eliminate "some types of violent behavior that cannot be controlled by any form of psychosocial therapy," said Samuel A. Corson, Ph.D., Professor of Psychiatry at Ohio State University College of Medicine.

Dr. Corson described the dog, Jackson, as spontaneously and aggressively vicious. A beagle-cocker spaniel hybrid, he responded to any approach with snapping, snarling, growling—or, if possible, biting—and in the course of a notorious career in the laboratory had attacked other dogs, bitten experienced and gentle handlers, and ruined considerable equipment when pavlovian conditioning was attempted.

Tranquillizers failed to help, and since

the 18-month-old dog also exhibited hyperkinesis the decision was made to try amphetamine, embedded for safety's sake in a meatball. The dosage approximated that used with hyperkinetic children.

Vicious barking and snarling disappeared

Continued on page 12

helps relieve anxiety-linked symptoms in duodenal ulcer

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Hyperkinetic dog before, I, and after d-amphetamine therapy, with Dr. Corson.

# Medical Tribune

and Medical News

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world news of medicine and its practice—fast, accurate, complete

Wednesday, February 14, 1973

## Test Employing Cold Stimulus Shows Sclerosis

Medical Tribune Report

SAN JUAN, P.R.—Intensive computerized testing has confirmed the hypothesis that the cold-pressor test, developed in the 1930s for indicating prehypertensive states and later virtually abandoned, is effective as a screening test for arteriosclerosis.

Dr. Ignatios J. Voudoukis, chief of the hypertension section of the Hutzel Hospital Unit, Wayne State University School of Medicine, said here that "excessive acute blood pressure elevations (systolic and pulse pressure) precipitated by a cold stimulus should be considered as an indication of clinically significant atherosclerotic vascular disease rather than hypertension."

Speaking at the 19th Annual Meeting of the American College of Angiology, Dr. Voudoukis suggested that "any individual with exaggerated cold-pressor response should be further investigated for clinically significant vascular sclerosis."

Cold-pressor response was determined in 64 consecutive ambulatory patients of a predominantly hypertensive population seen in a solo private practice. They were divided into four groups—83 patients free of hypertension and arteriosclerosis, 66 with arteriosclerosis, 93 with hypertension, and 399 who had hypertension with superimposed arteriosclerosis.

The two latest M.D. groups to organize comprised house staff members at the Contra Costa County Hospital, Martinez, Calif., and the municipally operated Jersey City (N.J.) Medical Center.

The action taken by the Contra Costa physicians followed upon the merger of

Continued on page 27

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## Vascular Operation Tried Successfully In Sexual Impotence

Medical Tribune World Service

PRAGUE—Microvascular surgery to transplant a saphenous vein segment has been used successfully here in the treatment of selected cases of sexual impotence at the Institute of Clinical and Experimental Medicine.

The first case was that of an automobile accident victim with pelvic fracture and extensive hematomata and internal bleeding in the pelvic and genital region, which required tying off of the internal iliac branches.

The patient was rendered impotent, and Dr. Vaslav Michal was asked to perform aortographic studies. These showed poor circulation to the entire pelvic region. Dr. Michal conducted a literature study, with meager results, he related: combinations of atherosomatous plaques and poor circulation in the lower extremities with impotence were known, but surgical attempts at correction were few and of doubtful value.

### Endarterectomy Considered

Several previous reports were concerned with iliac endarterectomy to improve circulation to the penis, but while 30 per cent of the patients showed some improvement in erection, another 30 per cent showed no change, and even in the improved cases, ejaculation had usually disappeared completely.

Dr. Michal believed that the latter complication came about because the surgery was intrapelvic and required interruption of the pelvic autonomic nerve plexuses involved in the ejaculation reflex. In his own first case, further intrapelvic surgery was out of the question, he said, since previous surgery had left the terrain unrecognized.

### Motorbike Mishaps Cited

Medical Tribune World Service

THE HAGUE—Young persons between the ages of 15 and 19 have a higher rate of hospitalization and mortality from accidents than any other age group in the Netherlands, according to a report by the Dutch Medical Registration Foundation.

Traffic accidents account for most of the cases, and most of the victims are riders of light motorcycles.

## Australians Claim Success in Program To Return Women Doctors to Medicine

Medical Tribune World Service

SYDNEY, AUSTRALIA—A plan to help women doctors return to the profession if they have been away from medicine for some time is working successfully here.

The three-month retraining course, instituted last year, is conducted by the Mater Misericordiae Hospital, North Sydney, under the guidance of the clinical superintendent, Dr. Geoffrey Dlethelm.

Five women have already completed the

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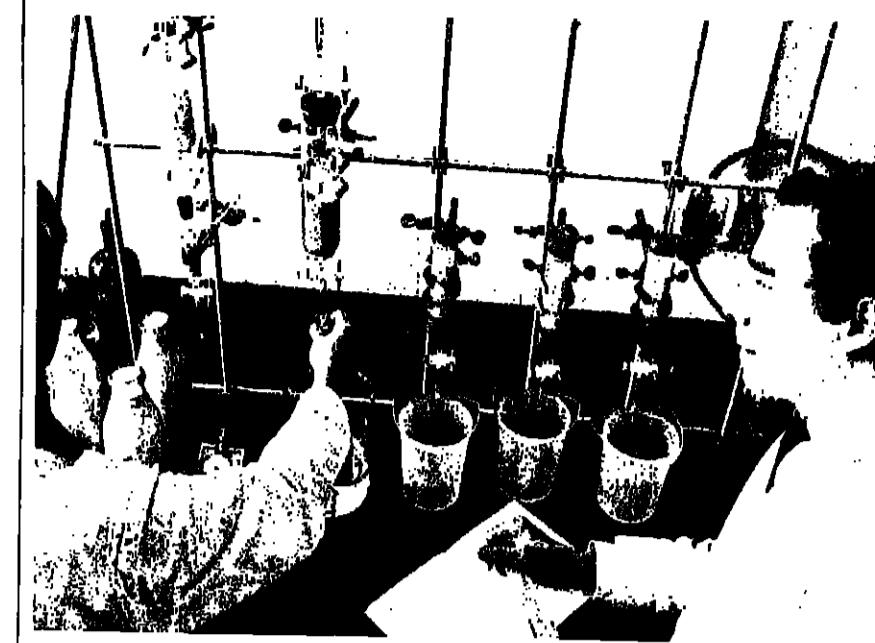
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## Importance of the Public Health Lab



The public health laboratory is an essential tool of every public health service in the world. The laboratory is needed to define the magnitude of certain disease problems, to determine the control strategy, and to help in appraising the degree of success in disease control. Above, at Chilean public health lab, milk is tested for strontium-90.

nizable, and so he began to work out an extrapelvic approach. This called for surgery carried out under a dissecting microscope with special instruments—a technique in which he had been trained during a year's stay with Prof. Julius H. Jacobson II at the Mount Sinai Hospital, New York.

The first approach tried, which worked completely and immediately, Dr. Michal reported, was to use a deep saphenous vein segment as a graft, attaching one end to the pudendal artery exposed from the perineum and other end to the medial side of the femoral artery, both junctions end-to-side, with the graft being led subcutaneously along the scrotum and then by tunneling into the femoral triangle. The microsurgery was necessitated by the small size of the graft and the pudendal artery. Sexual competence returned within a few days of surgery.

The operation itself, Dr. Michal commented, is simple, rapid, and relatively traumatic—two small incisions and only subcutaneous dissection. He performed the operation eight times on cadavers before the first clinical attempt.

After the first successful experience, he turned his attention to the far more common case of impotence caused by atherosomatous plaques, and developed an aortographic technique in order to analyze the vascular situation.

One of his main research interests at present is the development of a reliable diagnostic test for a vascular basis of impotence. His approach is to measure blood flow in the penis with either thermistor or impedance plethysmography. His problem is how to induce erection by constant and reliable technique, and he is trying to use such peptide drugs as vasopressin.

### Australian MD Group Is Opposed To New Government Health Plan

Medical Tribune World Service

CANBERRA—Health care plans by Australia's newly elected Labor Government face stiff opposition from the Australian Medical Association.

Timing for the introduction of Labor's proposed single-fund insurance plan—to be financed by a tax surcharge—will depend on the cooperation of the doctors, Prime Minister Edward Gough Whitlam has declared.

But the medical association has already announced it will oppose any move by the Government to abandon the present voluntary health insurance scheme or to turn physicians into salaried civil servants.

For the past three years the number of cases has not gone above what Dr. Bourgasov calls a "normal level" for influenza, but this year much of the population had lost the two or three years' immunity carried from the last attack of the virus.

"We haven't beaten the virus," Dr. Bourgasov commented, "but with the precautions we took we have been able to limit the spread of infection and to prevent in many cases the complications that increase mortality."

### Eradication of Smallpox By 1975 Is Foreseen

Medical Tribune World Service

GENEVA, SWITZERLAND—The World Health Organization predicted here that smallpox would be eradicated by 1975 if present programs are maintained. WHO's Executive Board reported that, although smallpox incidence last year increased to about 65,000 cases, that figure represented better reporting and diagnosis.

The health minister of that state, Dharan Datta Valdya, told a conference of eye surgeons in Patras that blindness caused by infectious diseases is on the decline, but that blindness caused by malnutrition is on the increase, particularly among children and expectant mothers.

With the exception of Bangladesh, Pakistan, and India, where major outbreaks occurred, there were relatively few cases in the world.

CLINICAL NEWS NOTE: "Insofar as it is valid to extrapolate from animals to humans, what hyperkinetic or violent children learn in school while medicated with amphetamines they would tend to retain later." (S. A. Corson, Ph.D., see page 1.)

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## Flu Epidemic in U.S.S.R. Cripples Schools, Stores And Taxes Health Service

Medical Tribune World Service

MOSCOW—Health services in the Soviet Union were emerging late last month from a battle on a massive scale with the A England 42-72 influenza virus.

At the peak of the epidemic, Moscow was reporting 70,000 new cases a day and Leningrad 30,000 a day. Computer tracking indicated that a second onslaught by the virus might be on the way.

Many schools were closed and subway services were reduced, even in the rush hours. Customers and salesgirls in the shops wore face masks. Production in factories and work in offices slowed to a snail's pace because of absenteeism.

Said one office manager: "Two or three people out with the influenza at this time of year is normal for us, but this season it's 15 or 20 at once. A lot of the work has just come to a standstill."

When Leningrad was alerted the virus was approaching, children were on year-end vacation, so the vacation was prolonged while certain day nurseries and crèches began working round the clock.

### Museum Queues Disappeared

The always familiar long queues of Moscow schoolchildren going to the museums disappeared during the epidemic. For those schools that were open, entries were banned.

In cinemas in Moscow, Leningrad, and other main cities, long intermissions were introduced between showings to allow disinfection of the premises.

An "influenza task force," headed by Dr. Piotr Bourgasov, Deputy Minister of Health, with a flu warning system linking 122 cities, was established. Its object was to advise the population on precautions to be taken and to organize and direct all health services mobilized for the battle.

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### Half of Blind Are Indian

Medical Tribune World Service

NEW DELHI—Physicians here estimate that India's blind now number some 10,000,000 persons, half the world total. Most of them live in Uttar Pradesh.

The health minister of that state, Dharan Datta Valdya, told a conference of eye surgeons in Patras that blindness caused by infectious diseases is on the decline, but that blindness caused by malnutrition is on the increase, particularly among children and expectant mothers.

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### ECTOPIC BEAT

"The low price includes round-trip jet transportation including meals and beverages; a double room with private balcony; American breakfasts every morning; five full-course dinners, including a Caribbean luau and a barbecue...."

—*Bulletin of the Beaver County (Pa.) Medical Society*

A Caribbean luau is a calypso hula, but what's a barbecue?

(Regular beat: Immunotherapy, page 35.)

## Child Health Center Serves Chicagoans



The Woodlawn Child Health Center, located on Chicago's South Side, provides free comprehensive health services to children in the Woodlawn area who need primary health care and preventive services. Above, Dr. Alberto Gedlman, one of four pediatricians supplied by the University of Chicago Pritzker School of Medicine, with patient. At right, Veronica Chandler works with the files of the Center's 16,000 registered patients.

Lab technician Beatrice Cain at work in the center's laboratory. According to Dr. John Madden, medical director, the presence of the center in the community has been a contributing factor to the improved general health around Woodlawn.

### Rheumatoid Arthritis

## Subgroup of Patients Responds to Histidine

Medical Tribune Report

PITTSBURGH—A study of histidine treatment in patients with rheumatoid arthritis suggests that a subgroup of patients with severe active disease of long duration may experience a modest degree of clinical improvement after having undergone this form of therapy.

This finding was reported at the interim scientific session of the American Rheumatism Association by Dr. Robert S. Pinals, of the State University of New York, Upstate Medical Center, Syracuse.

Physician evaluations were found to yield a somewhat similar pattern of results.

Because of the small group of patients studied and the variation in histidine levels on different days, it was not possible to make significant clinical correlations, said Dr. Pinals, but there was a suggestion that lower serum levels of histidine were associated with superior therapeutic results.

Levels increased in the histidine group but not in the placebo group during the study.

"On the basis of this study, we must say that the therapeutic efficacy of histidine has not yet been established and that general use of this treatment cannot be recommended," said Dr. Pinals.

### Response to Treatment Compared

Evaluation of response to treatment revealed no significant differences between the two groups in grip strength, sedimentation rate, walking time, morning stiffness, or number of swollen and tender joints, Dr. Pinals reported.

Neither was there significant improvement in these parameters within each group, except in hematoctrit in the histidine-treated patients and the grip strength in the placebo group.

The selections were announced by Dr. Arthur James, president of the A.C.S., and Dr. Frank J. Rauscher, Jr., director of the NCI.

The three sites are: the Stella and Charles Guttman Breast Diagnostic In-

### Imprinters to Help MDs Cut Misuse of Drugs

Medical Tribune Report

CINCINNATI—To help pharmacists in dispensing medication and to reduce prescription forgeries, the University of Cincinnati Medical Center will issue imprinters to physicians on the house staff. The imprinter is a stamp with the physician's name and identification number.

It is also hoped that they will eliminate inconvenience experienced by patients when prescriptions cannot be filled because of illegible signatures.

This program to control hospital prescription blanks is the first in Ohio and may be unique in the nation, the medical center said.

The innovation was suggested by Robert Bundman, chief pharmacist at Holmes Hospital.

Co-workers were Drs. Edward D. Harris, Jr., and James Frizzell, of Dartmouth Medical School and Dr. Donald A. Gerber, of the State University of New York, Downstate Medical Center, Brooklyn.

Mr. Chase commented: "I'm especially pleased that we have been able to get the program off the ground so fast. So far as I know, no other government in the country is doing anything like the massive testing we are doing."

"HSA's screening program already has been very useful in term of public education. We've helped to make more New Yorkers aware that high blood pressure is a serious health problem.

"Now that HSA's program has shown many New Yorkers that they have high blood pressure, we are concerned about what these people do with that information. We suspect that many do nothing more than make a mental note of it. The major thrust of HSA planning for hypertension control in 1973 will be to set up and evaluate pilot treatment programs for victims of high blood pressure."

## Exposure to Cadmium May Pose Threat to Man

Medical Tribune Report

WEST LAFAYETTE, IND.—Exposure to cadmium may pose an environmental threat to man, a team of 16 Purdue University students reported to the National Science Foundation.

The team spent 11 weeks last summer investigating the levels of cadmium in the environment in a program called Student-Originated Studies, cosponsored by the National Science Foundation and Purdue's Institute for Environmental Health. The director of the Purdue project was John E. Christian, Ph.D., chairman of the Department of Bionucleonics.

Utilizing radioisotopes and radiation counters in one portion of the investigation, the students found that all species studied exhibited high retention of cadmium each day. Although the dispersion of cadmium in food chains is poorly monitored, and concentrations in normal diets must necessarily be approximated, it is estimated that the daily oral human in-

take in industrial areas lies in the range of 200 and 400 micrograms."

# natural superiority



Naturally, an imitation does not equal the original. Synthetic chemicals often lack some vital factors present in the natural medicinal.

Take SENOKOT Tablets/Granules, for example. This highly effective laxative gets a head start from Mother Nature—natural senna from the *Cassia acutifolia* plant has been used as a laxative for over 1500 years. In SENOKOT® preparations, this natural vegetable laxative is purified and refined into one of the most modern, virtually colon-specific, predictably gentle anticonstipants your patients' care.

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**Senokot**  
TABLETS GRANULES  
(standard senna concentrate)  
A natural laxative

## WHAT'S NEW IN RHEUMATOLOGY

### What's new and important in rheumatology?—I

#### The Consultant

DR. LEE E. BARTHOLOMEW

Professor of Rheumatology,  
Head, Division of Rheumatology,  
Albany Medical College, Union University, Albany, N. Y.



THE IMMUNOLOGY of connective-tissue diseases probably takes the forefront at the present time. Systemic lupus erythematosus, being the prototype of the immune complex diseases, is the subject of much interesting and exciting work. The fact that there are several antinuclear antibodies which have been described, and probably many more yet to come, provides considerable interest to the possibilities

The basic conservative program of the treatment of rheumatoid arthritis in the adult consists of adequate salicylates; that is, blood levels between 15 and 25 mg. per 100 ml. two or three hours after tak-

ing their last dose of aspirin, adequate rest (both body rest and joint rest) for the acute phases of the disease. Simple measures, such as cock-up splints for the hands and wrists to be worn at night and during the day, are extremely useful during acute flares of synovitis involving those joints. The third basic conservative measure is that of physical therapy, which includes not only the use of heat, such as the Hubbard tank, paraffin to the hands and fingers, hot packs, Hydrocollator packs, etc., but also the cautious use of range-of-motion exercises to prevent deformities and muscle-strengthening exercises of individual muscle groups which have become atrophied.

This basic program is given for periods of two to four months. If at the end of that time there has not been adequate suppression of this disease in terms of decrease in morning stiffness, fewer joints showing active synovitis, increase in well-being and less general fatigue and malaise, and improvement in sedimentation rate and anemia, additional therapy is then indicated. I personally feel that the use of intramuscular gold salts is not only the most potent but the most effective of the anti-inflammatory agents used for rheumatoid arthritis. It is not without its hazards and toxicity, and for this reason great care should be taken in using the gold salts.

This requires a cooperative patient, a patient who is willing to come in to see the physician regularly. Before each injection, complete blood counts, evaluation of platelets, complete urinalysis are performed, the patient is observed for skin rashes, oral mucous membrane lesions, and questioned concerning whether they are developing any pruritis.

Patients are usually given 5 to 10 mg. at the first injection, 25 at the second, and then 50 mg. weekly until approximately 1 Gm. of gold has been given. Usually, if response is to occur it begins somewhere between 500 and 1,000 mg., and if they respond well, a maintenance program is established for an indefinite period using 50 mg. intramuscularly every month. If signs of toxicity occur the drug is withheld, or if significant toxicity occurs the drug is stopped completely. Using this cautious approach, rarely do significant toxic reactions occur.

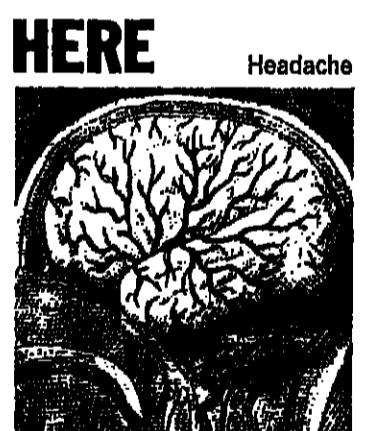
In general, one can expect that approximately 50-60 per cent of patients who can tolerate the drug will have improvement. Many of them will have a complete remission that may last for years.

Next week Dr. Bartholomew will discuss the immunologic aspects and treatment of systemic lupus erythematosus.

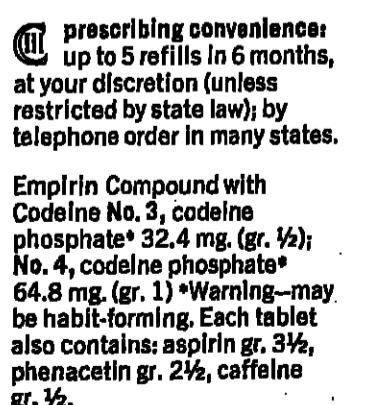
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Wherever it hurts, Empirin Compound with Codeine usually provides the symptomatic relief needed.



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#3, codeine phosphate\* (32.4 mg.) gr. 1/2  
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# Questions doctors are asking about Tranxene® (CLORAZEPATE DIPOTASSIUM)

4306 CB



## the new benzodiazepine from Abbott

### About these questions, Doctor:

Since the introduction of our new anti-anxiety agent, Tranxene (clorazepate dipotassium), in early October, we have maintained an 8:00-to-5:00, private-line communication system with our field representatives for the purpose of gathering and answering questions being raised daily by physicians.

The questions presented here are among those most frequently discussed, and the answers reflect the best available information to date.

Perhaps you'll find questions of your own here. In any event, we hope you'll find them useful.

#### Q. What is the fate of the drug in the body?

A. The drug is metabolized in the liver and excreted primarily in the urine.

#### Q. Does drug accumulation occur?

A. When recommended daily doses are administered, drug accumulation in the serum occurs only up to the seventh day. At this time, a plateau is reached and serum levels tend to remain stable with continued administration of the original dose.

#### Q. What is the half-life of Tranxene?

A. The serum half-life of nordiazepam, the primary metabolite of Tranxene, is approximately one day.

#### Q. What is the oral LD<sub>50</sub>?

#### Q. What is the effect on blood pressure?

A. Decreases in systolic blood pressure have been observed. In our premarketing clinical studies, the only effect seen on blood pressure was the lowering of slightly elevated systolic blood pressure in some patients.

#### Q. Does Tranxene cause bradycardia?

A. There were no reports of bradycardia in the controlled premarketing clinical studies on Tranxene.

#### Q. Can urinary retention be associated with Tranxene?

A. Anti-cholinergic effects have been reported with some benzodiazepines, and therefore, it may be possible that these effects could be seen with Tranxene as well.

#### Q. What is the rate of excretion?

A. After a single dose, approximately fifty percent is excreted primarily in the urine in the first 24 hours. By the tenth day, 80 percent of the drug is excreted. At that point, the excretion rate was found to be about one percent per day.

#### Q. Has respiratory depression been seen in the studies with Tranxene?

A. There was no evidence from our premarketing clinical studies demonstrating respiratory depression with the use of recommended doses of Tranxene. However, since it is a CNS depressant, one can assume that if massive doses were ingested, respiratory depression could occur.

#### Q. Does Tranxene affect the SGOT level?

A. In the clinical studies, there were reports of occasional increases of SGOT level in some patients. Increases of SGOT level have been reported with other benzodiazepines.

#### Q. Does this mean that Tranxene is contraindicated for anyone with impaired liver function?

A. It is not a contraindication. However, as with all benzodiazepines, the usual precautions in treating patients with impaired liver function should be observed.

#### Q. What is the oral LD<sub>50</sub>?

A. In rats the LD<sub>50</sub> was 1320 mg./kg; in monkeys the LD<sub>50</sub> could not be determined because of the emetic effect of large doses, but the LD<sub>50</sub> exceeds 1600 mg./kg.

#### Q. Is it true that Tranxene can cause a decrease in hematocrit?

A. Decreases in hematocrit have been reported. A causal relationship has not been established.

#### Q. Can the actions of Tranxene be potentiated by the concurrent use of other drugs? What about sedation?

A. Like other benzodiazepines, the actions of Tranxene may be potentiated by the concurrent use of barbiturates, narcotics, phenothiazines, monoamine oxidase inhibitors or other antidepressants. Clinical studies have shown increased sedation with concurrent use of hypnotics.

#### Q. Does Tranxene have muscle relaxant properties?

A. Clinical studies in muscle relaxation have not been performed.

#### Q. If Tranxene is administered to elderly patients with symptoms of anxiety, what special precautions should be observed?

A. An important precaution which should be taken when prescribing Tranxene for an elderly patient is to follow the patient closely at the initiation of therapy to observe his response. In elderly or debilitated patients, it is advisable to initiate therapy at a daily dose of 7.5 mg. to 15 mg., rather than the usual recommended daily dose of 30 mg.

Therapy should take into account possible drug interactions since the elderly patient may be on other drugs.

#### Q. How long was Tranxene studied before being introduced?

A. The clinical investigation of Tranxene was conducted for over four years in the United States. The investigation included studies ranging from three weeks to six months.

### Is Tranxene® effective? (CLORAZEPATE DIPOTASSIUM)

#### Physician Evaluations:

In double-blind clinical studies, Tranxene was shown to be effective in relieving symptoms of anxiety.

#### Patient Evaluations:

In most clinical studies, a series of patient self-evaluation tests were conducted under double-blind conditions before, during and after study. Improvement was recorded as a reduction in number or severity of anxiety symptoms.

Patient self-evaluations correlated well with physician evaluations—i.e. patients rated most improved by physicians tended to show greatest reduction in symptom test scores.

By both physician and patient assessment, therapy with Tranxene had a measurable effect in reducing the number and severity of symptoms.

#### Tranxene® is provided in 3 strengths: CLORAZEPATE DIPOTASSIUM



Tranxene is administered orally in divided doses; usual daily dose is 30 mg. The dose should be adjusted gradually within the range of 15 to 60 mg., based on response of the patient. In elderly or debilitated patients, it is advisable to initiate therapy at a daily dose of 7.5 mg. to 15 mg.

**In the management of anxiety... If you measure the success of the therapy by the patient's response, Tranxene (CLORAZEPATE DIPOTASSIUM) is an effective measure.**

See last page for prescribing information.



# In the management of anxiety... If you measure the success of the therapy by the patient's response,

## Tranxene® is an effective measure. (CLORAZEPATE DIPOTASSIUM)

### Tranxene® (CLORAZEPATE DIPOTASSIUM)

**DESCRIPTION:** Chemically, TRANXENE (clorazepate dipotassium) is a benzodiazepine. The empirical formula is  $C_{18}H_{11}ClK_2N_2O_4$ ; the molecular weight is 408.93. The compound occurs as a fine, light yellow, practically odorless powder. It is insoluble in the common organic solvents, but very soluble in water. Aqueous solutions are unstable, clear, light yellow, and alkaline.

**ACTIONS:** Pharmacologically, TRANXENE (clorazepate dipotassium) has the characteristics of the benzodiazepines. It has depressant effects on the central nervous system. The primary metabolite, nordiazepam, reaches peak level in the blood stream at approximately 1 hour. The plasma half-life is about 1 day. The drug is metabolized in the liver and excreted primarily in the urine. (See ANIMAL AND CLINICAL PHARMACOLOGY section).

**INDICATIONS:** TRANXENE is indicated for the symptomatic relief of anxiety associated with anxiety neurosis, in other psychoneuroses in which anxiety symptoms are prominent features, and as an adjunct in disease states in which anxiety is manifested.

**CONTRAINDICATIONS:** TRANXENE (clorazepate dipotassium) is contraindicated in patients with a known hypersensitivity to the drug, and in those with acute narrow angle glaucoma.

**WARNINGS:** TRANXENE is not recommended for use in depressive neuroses or in psychotic reactions.

Patients on TRANXENE should be cautioned against engaging in hazardous occupations requiring mental alertness, such as operating dangerous machinery including motor vehicles.

Since TRANXENE has a central nervous system depressant effect, patients should be advised against the simultaneous use of other CNS-depressant drugs, and cautioned that the effects of alcohol may be increased.

Because of the lack of sufficient clinical experience, TRANXENE (clorazepate dipotassium) is not recommended for use in patients less than 18 years of age.

**Physical and Psychological Dependence:** Withdrawal symptoms (similar in character to those noted with barbiturates and alcohol) have occurred following abrupt discontinuance of clorazepate. Symptoms of nervousness, insomnia, irritability, diarrhea, muscle aches and memory impairment have followed abrupt withdrawal after long-term use of high dosage.

Caution should be observed in patients who are considered to have a psychological potential for drug dependence.

Evidence of drug dependence has been observed in dogs and rabbits which was characterized by convulsive seizures when the drug was abruptly withdrawn or the dose was reduced; the syndrome in dogs could be abolished by administration of clorazepate.

**Usage in Pregnancy:** Reproduction studies have been performed in rats and rabbits and there was no evidence of harm to the animal fetus. The relevance to the human is not known. Since there is no experience in pregnant women who have received this drug, safety in pregnancy has not been established.

It is assumed that TRANXENE or its metabolites is

excreted in human milk. Therefore, this drug should not be given to nursing mothers.

**PRECAUTIONS:** In those patients in which a degree of depression accompanies the anxiety, suicidal tendencies may be present and protective measures may be required.

The least amount of drug that is feasible should be available to the patient.

Patients on TRANXENE for prolonged periods should have blood counts and liver function tests periodically. The usual precautions in treating patients with impaired renal or hepatic function should also be observed.

In elderly or debilitated patients, the initial dose should be small, and increments should be made gradually, in accordance with the response of the patient, to preclude ataxia or excessive sedation.

**ADVERSE REACTIONS:** The side effect most frequently reported was drowsiness. Less commonly reported (in descending order of occurrence) were: dizziness, various gastrointestinal complaints, nervousness, blurred vision, dry mouth, headache, and mental confusion. Other side effects included insomnia, transient skin rashes, fatigue, ataxia, genito-urinary complaints, irritability, diplopia, depression and slurred speech.

There have been reports of abnormal liver and kidney function tests and of decrease in hematocrit.

Decrease in systolic blood pressure has been observed.

**DOSAGE AND ADMINISTRATION:** TRANXENE (clorazepate dipotassium) is administered orally in divided doses. The usual daily dose is 30 mg. The dose should be adjusted gradually within the range of 15 to 60 mg. daily in accordance with the response of the patient. Drowsiness may occur at the initiation of treatment and with dosage increments. In elderly or debilitated patients it is advisable to initiate treatment at a daily dose of 7.5 to 15 mg.

**DRUG INTERACTIONS:** If TRANXENE (clorazepate dipotassium) is to be combined with other drugs acting on the central nervous system, careful consideration should be given to the pharmacology of the agents to be employed. Animal experience indicates that TRANXENE prolongs the sleeping time after hexobarbital or after ethyl alcohol, increases the inhibitory effects of chlorpromazine, but does not exhibit monoamine oxidase inhibition. Clinical studies have shown increased sedation with concurrent hypnotic medications. The actions of the benzodiazepines may be potentiated by barbiturates, narcotics, phenothiazines, monoamine oxidase inhibitors or other anti-depressants.

If TRANXENE is used to treat anxiety associated with somatic disease states, careful attention must be paid to possible drug interaction with concomitant medication.

**MANAGEMENT OF OVERDOSE:** As in the management of overdose with any drug, it should be borne in mind that multiple agents may have been taken.

If vomiting has not occurred spontaneously, it should be induced. Immediate gastric lavage is also recommended. General supportive care, including frequent monitoring of the vital signs and close observation of the patient, is indicated.

Hypotension, though unlikely, may be controlled with Levophed® (levarterenol) or Aramine® (metaraminol). Caffeine and Sodium Benzoate Injection, U.S.P., may be used to counteract central nervous system depressant effects.

There has been reported a 41-year-old woman who took 25 capsules (187.5 mg.) of TRANXENE. Severe diarrhea and vomiting occurred, but she made an uneventful recovery without being hospitalized.

**ANIMAL AND CLINICAL PHARMACOLOGY:** Studies in rats and monkeys have shown a substantial difference between doses producing tranquilizing, sedative and toxic effects. In rats, conditioned avoidance response was inhibited at an oral dose of 10 mg./kg.; sedation was induced at 32 mg./kg.; the LD<sub>50</sub> was 1320 mg./kg. In monkeys aggressive behavior was reduced at an oral dose of 0.25 mg./kg.; sedation (ataxia) was induced at 7.5 mg./kg.; the LD<sub>50</sub> could not be determined because of the emetic effect of large doses, but the LD<sub>50</sub> exceeds 1600 mg./kg.

Twenty-four dogs were given TRANXENE orally in a 22-month toxicity study; doses up to 75 mg./kg. were given. Drug-related changes occurred in the liver: weight was increased and cholestasis with minimal hepatocellular damage was found, but lobular architecture remained well preserved.

Eighteen rhesus monkeys were given oral doses of TRANXENE from 3 to 36 mg./kg. daily for 52 weeks. All treated animals remained similar to control animals. Although total leucocyte count remained within normal limits it tended to fall in the female animals on the highest doses.

Examination of all organs revealed no alterations attributable to TRANXENE. There was no damage to liver function or structure.

**Reproduction Studies:** Standard studies of fertility, teratology and reproduction were conducted on rats and rabbits. Oral doses in rats up to 150 mg./kg. and in rabbits up to 15 mg./kg. produced no abnormalities in the fetuses and no impairment to fertility and reproductive capacity of adult animals attributable to TRANXENE (clorazepate dipotassium). As expected, the sedative effect of high doses interfered with care of the young by their mothers (see Use in Pregnancy).

**Clinical Pharmacology:** Studies in healthy men have shown that TRANXENE has depressant effects on the central nervous system. Prolonged administration of high doses (120 mg. daily as a single oral dose) was without toxic effects, and abrupt cessation of drug was not followed by serious signs or symptoms.

**Absorption—Excretion:** After oral administration of TRANXENE (clorazepate dipotassium), there is essentially no circulating parent drug. Nordiazepam, its primary metabolite, quickly appears in the blood stream with peak levels at about 1 hour. The plasma half-life is approximately 1 day. In volunteers given 15 mg. (50 μC) of <sup>14</sup>C-Tranxene, about 80% was recovered in the urine and feces within 10 days. Excretion was primarily in the urine with about 1% excreted per day on day 10.

**HOW SUPPLIED:** TRANXENE (clorazepate dipotassium) is supplied as capsules, in bottles of 100.

The capsules contain:

3.75 mg (gray with white cap).....	NDC 074-3417-13
7.5 mg (gray with maroon cap).....	NDC 074-3418-13
15 mg (all gray).....	NDC 074-3419-13

## Doctors' Debate

MEDICAL TRIBUNE frequently receives extensive and well-documented communications from physicians on current subjects of controversy or those of great current medical interest. We invite contributions in these areas for presentation in this new feature.

In the January 10 issue of MEDICAL TRIBUNE, Dr. Seymour Diamond, Assistant Professor of Neurology at the Chicago Medical School and president of the American Association for the Study of Headache, was "In Consultation" (page 5) on the subject: What's new and important in headache study? Dr. Warren F. Wilhelm, of Kansas City, Mo., then submitted a question to Dr. Diamond in a letter to the editor. Following is the question and Dr. Diamond's answer:

### QUESTION

What questions should elicit a reasonable and thorough headache history?

### ANSWER

We carefully question all patients regarding the onset of their headache symptoms. Whether headache occurred for the first time in childhood or late in life can sometimes predetermine what type of headache it is. Most migraine headaches will appear in childhood or teens and be present at least through the 50s. Headaches due to depression occur most commonly in the 40-60 age group but can occur at any age.

**Location of headache:** Most migraine headaches and cephalgias due to organic disease are one-sided, while headaches due to psychogenic causes are generalized, having a hatbandlike distribution.

**Frequency:** A headache occurring every day most often is psychogenic, but certain persistent migraine headaches and cluster headaches can occur daily. A brain tumor will give an unrelenting headache.

**Duration:** A headache that is constant and never relents is most often psychogenic or due to organic disease.

**Severity:** Sometimes a clue because a headache due to psychogenic causes is not very severe, while migraine headache has a moderate to great severity and in cluster headache the pain is sometimes so great as to make the patient want to commit suicide.

It has been suggested that I should seek to win over the medical establishment in this country. I have been in constant communication with, and have been scorned by, nearly all our ob/gyn authorities, by our "nutrition experts," pediatricians, nursing authorities, pathologists, and journal editors, especially the *New England Journal of Medicine*, the *American Journal of Obstetrics & Gynecology*, *SG&O*, and *OB/GYN Survey*.

Private pharmaceutical companies are no better, as they continue to push diuretics, appetite depressants, and salt substitutes for use in pregnancy. For over six years I have had a constant battle against these practices. Worse of all are the Federal and state bureaus and institutes charged with protecting the public health, including HEW, FDA, and the USPHS.

I and others have published a wide range of statistics and many clinical studies to prove the importance of good nutrition—and the dangers of weight restriction, salt restriction, and salt diuretics for gravid women. There is an extensive and sound medical literature on this subject, available to those who wish it.

Perhaps, instead of cold statistics, a case history may make the point more vividly:

Patient M. was a small Mexican woman who followed her doctor's orders to the letter. A private ob/gyn specialist in California restricted her to one egg and one glass of milk a week, on the grounds that there is too much salt in milk and eggs. She was constantly advised at each prenatal visit: "Keep your weight down! Keep your weight down!" She wanted a healthy baby, so she faithfully followed her doctor's orders. Result: she gained only 14 pounds in all (from 112 to 126) and went into labor right at term. This was three months after she had been given a low-salt diet and diuretic pill to take every day; she didn't miss a day.

Her son, J.F., weighed 4 pounds, 15 ounces at birth. His blood sugar dropped to 20 mg. per cent and then later to 12 mg. per cent, and he had hypoglycemic convulsions repeatedly. The mother, after a normal blood loss at delivery, went into what her doctor termed "idiopathic shock"—which we know was caused by her hypovolemia.

The boy is obviously and grossly mentally retarded and has to attend a special school for brain-damaged children. At age 15 months he was age three to four months in development and function on the Denver Grid—head drop, crossed eyes, small head. At age 18 months he still could not pull to stand or walk.

The patient had her second son after prenatal care in my clinic. During this second pregnancy she gained 50 pounds, had two eggs and a quart of milk every day, meat, vegetables, fruits, cereals, and no salt diuretics, no dietary salt restriction. She was told on each visit: "Keep eating a good diet—salt your food to taste!" This second child, A., weighed 9 pounds at birth and is a perfect specimen.

Fellow American physicians, how long are we going to disregard the scientific evidence of the causal relationship of protein-calorie malnutrition, restriction of salt, and the dangerous use of salt diuretics to complications of pregnancy, fetal mortality, and damage to the newborn human infant?

**Tom Brewer, M.D.**  
County Physician  
Richmond Health Clinic  
Richmond, Calif.

and moderate exercise in the treatment of consumption (tuberculosis), dropsy (heart-failure), and hypochondriacal dis temper (possibly manic-depressive psychosis).

I have completed the sentence which was bifurcated by your editorialist so that Fuller's true sentiment is expressed, as follows:

"That the Use of Exercise does conduct very much to the Preservation of Health, that it promotes the Digestions, rinses the Spirits, refreshes the Mind, and that it strengthens and relieves the whole Man, is scarcely disputed by any; but that it should prove Curative in some particular Distempers, and that too when scarce anything else will prevail, seems to obtain little credit with most People, who tho' they will give a Physician the hearing, when he recommends the frequent use of Riding, or any other sort of Exercise; yet at the bottom look upon it as a forlorn Method, and the Effects rather of his disability to relieve 'em, than of his Belief that there is any great matter in what he advises: *Thus by a negligent Indifference they deceive themselves, and let slip the Golden Opportunities of recovering, by a diligent Struggle, what could not be procured by the use of Medicine alone*" (italics mine).

**Allan J. Ryan, M.D.**  
Acting Chairman  
Committee on Exercise  
and Physical Fitness, A.M.A.

## Student, Teacher: Electronics Aids In Communication

**Medical Tribune Report**  
Los ANGELES—An \$80,000 electronic student response system, designed to increase the efficiency of student-teacher communication, is in operation at the University of Southern California School of Medicine.

The system, recently installed in the Louis B. Mayer Medical Teaching Center, allows individual student participation and response, which would otherwise be impossible in the large-classroom environment of the 500-seat auditorium.

As questions are presented by the Instructor, a push-button device on the arm of 265 seats allows a student to pick one of five possible answers. The device immediately indicates to the student whether he is right or wrong, and indicates to the Instructor the percentage of the class responding, and percentage correct or incorrect for each possible answer.

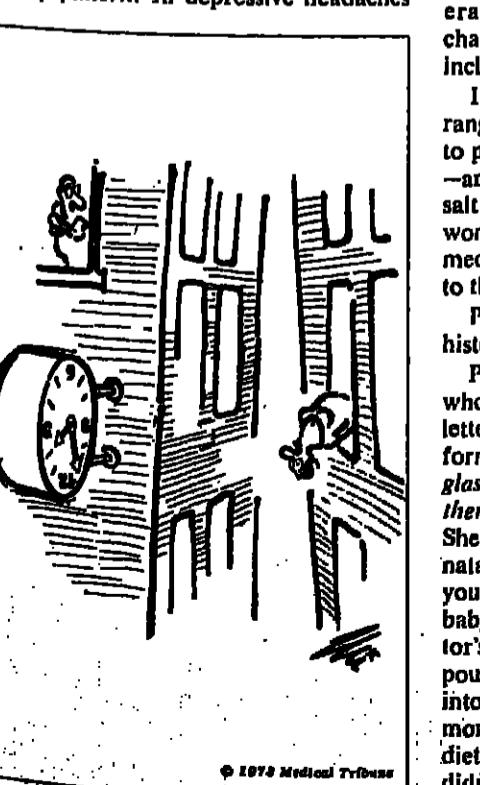
An electronic scanner collects the individual student responses and feeds them to a computer, which analyzes the data and relays it to a teletype. The Instructor receives an immediate printed readout with detailed data analysis of question-by-question performance by individual students and the class as a whole.

Thus, the Instructor can rapidly assess student understanding of materials presented, and identify areas that need reinforcing.

This system was described as representing a marked advantage over the traditional method of assessing student comprehension by giving quizzes, which have to be graded and then returned to the students—a process entailing a long interval between presentation of the material and determination of the extent of its assimilation.

As Dr. Phil Manning, Professor of Medicine and associate dean for postgraduate medical education, noted, "the new system will allow the U.S.C. faculty to organize problem-solving sessions with active participation in large groups. Those activities have previously been restricted to small groups."

The system was installed by Instructional Industries Inc., an independent affiliate of General Electric and an outgrowth of an educational systems group in the G.E. Research and Development Center.



# G.I. FORUM

A CURRENT REVIEW OF INVESTIGATIONS IN GASTROENTEROLOGY



## A kaleidoscopic entity

Gastritis...a disease of myriad uncertainties...a disease surrounded by much confusion. Very few subjects in medicine arouse so much difference of opinion.<sup>1</sup> Gastritis was discarded as a specific entity



## Gastroscopy alone or confirming biopsy?

Part of the confusion surrounding the diagnosis of gastritis lies in the difficulty of defining its various forms, which are largely determined by the diagnostic method used.<sup>2</sup> Gastroscopic definitions, based on direct visual inspection, do not always correlate well with the histologic state of the mucosa—which in turn may show little relationship to symptoms.<sup>3</sup> While some clinicians once considered gastroscopy to be the best method of diagnosing chronic gastritis,<sup>4</sup> most insist that the visual method be confirmed by biopsy.<sup>2,4</sup> The consensus is that despite the possibility of sampling error due to the limited area examined, histologic findings are the *sine qua non* in the classification of chronic gastritis.<sup>2</sup>

## Does aspirin irritate normal G.I. mucosa?

Almost always. Some view aspirin irritation of gastric mucosa as a general phenomenon rather than one restricted to hypersensitive persons.<sup>5</sup> Others suspect an individual sensitivity that develops only in particular circumstances.<sup>6</sup> Wide variations have been noted in individual tolerance of gastric mucosa to circulating salicylates.<sup>7</sup> One investigator suggests that those who are immune to aspirin irritation may have a high replacement of gastric epithelial cells.<sup>8</sup>

## Does gastritis precede ulcer or vice versa?

In more than 40 per cent of gastric ulcers, gastritis either appears as a border of swelling around the ulcer or involves all of the gastric mucosa.<sup>9</sup> But the

question of which came first—the ulcer or the gastritis—has never been settled. An old theory which still has its adherents regards the gastritis as secondary to the stomach ulcer.<sup>10</sup> This group saw it as an inflammatory reaction spreading from the ulcer site and usually called it "zonal gastritis." However, recent work using biopsy specimens obtained during gastroscopy would seem to refute this belief.<sup>10</sup> The persistence of superficial or atrophic gastritis after a gastric ulcer has healed would imply that the ulcer may be secondary to gastritis.

## The need to provide a comprehensive medical regimen

Such symptoms as anorexia, epigastric discomfort after meals, nausea, bloating and burning sensations may be sufficiently severe and persistent to require medical attention. Furthermore, if an acute stage of gastritis is left untreated, some clinicians feel that there is risk of its leading to chronic superficial gastritis, with possible progression toward gastric atrophy. Besides physical rest and respite for the inflamed stomach, some patients will very likely need respite from undue anxiety as well.

**References:** 1. Truelove, S. C., and Reynell, P. C.: *Diseases of the Digestive System*, Oxford, Blackwell Scientific Publications, 1963, p. 122. 2. Villardell, R.: "Chronic Gastritis" in Buckius, H. L.: *Gastroenterology*, ed. 2, Philadelphia, W. B. Saunders Co., 1963, vol. 1, pp. 368-404. 3. Schneider, E.: "Gastritis," in Pandorf, M. (ed.): *Gastroenterologic Medicine*, Philadelphia, Lea & Febiger, 1969, pp. 687-705. 4. Palmer, F. D.: *Clinical Gastroenterology*, ed. 2, New York, Hoeber Medical Division, Harper & Row, 1963, pp. 145-150. 5. Croft, D. N.: *Brit. Med. J.*, 2:164, 1961. 6. Lange, H. R.: *Gastroenterology*, 33:270, 1957. 7. Pen, D. J., and Wood, P. H.: *Gut*, 8:301, 1967. 8. Croll, D. N., Lame, 2:831, 1968. 9. Joske, R. A.; Finek, B. S., and Wood, I. J.: *Quart. J. Med. New Series*, 24:269, 1959. 10. Glear, M. W. I.; Truelove, S. C., and Whitehead, E. *Gut*, 12:639, 1971.

## The value of dual-action adjunctive therapy

For patients with acute gastritis who are experiencing both gastric distress and undue anxiety...Librax® is frequently useful adjunctive therapy. It provides the actions of both Librium® (chloridiazepoxide HCl) and Quarzan® (clidinium Br) in a single capsule that can help relieve the patient's excessive anxiety and provide antisecretory/antispasmodic action.

The value of Librium (chloridiazepoxide HCl) has been demonstrated whenever excessive anxiety or undue tension are significant components of the clinical profile. Experimental and clinical studies with Quarzan (clidinium Br) have shown that this agent exerts antisecretory and antispasmodic effects on the G.I. tract. These are two good reasons for you to prescribe adjunctive Librax as part of your medical regimen in treating gastritis.

## Up to 8 capsules daily in divided doses

For optimum response, dosage may be adjusted according to your patient's requirements, within the range of 1 or 2 capsules, 3 or 4 times daily.

Before prescribing, please consult complete product information, a summary of which follows.

**Indications:** Symptomatic relief of hypersecretion, hypermotility and anxiety and tension states associated with organic or functional gastrointestinal disorders; and as adjunctive therapy in the management of peptic ulcer, gastritis, duodenitis, irritable bowel syndrome, spastic colitis, and mild ulcerative colitis.

**Contraindications:** Patients with glaucoma; prostatic hypertrophy and benign bladder neck obstruction; known hypersensitivity to chloridiazepoxide hydrochloride and/or clidinium bromide.

**Warnings:** Caution patients about possible combined effect with alcohol and other CNS depressants. As with all CNS-acting drugs, caution patients against hazardous occupations requiring complete mental alertness (e.g., operating machinery, driving). Though physical and psychological dependence have rarely been reported on recommended doses, use caution in administering Librium (chloridiazepoxide hydrochloride) to known addiction-prone individuals or those who might increase dosage; withdrawal symptoms (including convulsions), following discontinuation of the drug and similar to those seen with barbiturates, have been reported. Use of any drug in pregnancy, lactation, or in women of childbearing age requires that its potential benefits be weighed against its possible hazards. As with all anticholinergic drugs, an inhibiting effect on lactation may occur.

**Precautions:** In elderly and debilitated, limit dosage to smallest effective amount to preclude development of ataxia, oversedation or confusion (not more than two capsules per day initially; increase gradually as needed and tolerated). Though generally not recommended, if combination therapy with other psychotropics seems indicated, carefully consider individual pharmacologic effects, particularly in use of potentially sedating drugs such as MAO inhibitors and phenothiazines. Observe usual precautions in presence of impaired renal or hepatic function. Paradoxical reactions (e.g., excitement, stimulation and acute rage) have been reported in psychiatric patients. Employ usual precautions in treatment of anxiety states with evidence of impending depression; suicidal tendencies may be present and protective measures necessary. Variable effects on blood coagulation have been reported very rarely in patients receiving the drug and oral anticoagulants; causal relationship has not been established clinically.

**Adverse Reactions:** No side effects or manifestations not seen with either compound alone have been reported.

with Librax. When chloridiazepoxide hydrochloride is used alone, drowsiness, dizziness and confusion may occur, especially in the elderly and debilitated. These are reversible in most instances by proper dosage adjustment, but are also occasionally observed at the lower dosage ranges. In a few instances syncope has been reported. Also encountered are isolated instances of skin eruptions, edema, minor menstrual irregularities, nausea and constipation, extrapyramidal symptoms, increased and decreased libido—all infrequent and generally controlled with dosage reduction; changes in EEG patterns (low-voltage fast activity) may appear during and after treatment; blood dyscrasias (including agranulocytosis) and jaundice and hepatic dysfunction have been reported occasionally with chloridiazepoxide hydrochloride; making periodic blood counts and liver function tests advisable during protracted therapy. Adverse effects reported with Librax are typical of anticholinergic agents, i.e., dryness of mouth, blurring of vision, urinary hesitancy and constipation. Constipation has occurred most often when Librax therapy is combined with other spasmolytics and/or low residue diets.

helps relieve anxiety-linked symptoms in gastritis

adjunctive  
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Each capsule contains 5 mg chloridiazepoxide HCl and 2.5 mg clidinium Br.

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## Haptene Said to Avert Allergy to Penicillin

Medical Tribune World Service

MONTRÉAL—Penicillin allergy can be averted by the monovalent haptene BPO-FLYS (benzylpenicilloylformyllysine), Dr. Alain de Weck, of the University of Berne, Switzerland, reported here.

Dr. de Weck, who is director of the institute of clinical immunology at the university, spoke at a conference on control of reagent-mediated hypersensitivity.

In clinical trials, he said, allergic reaction to penicillin could be prevented in 12 out of 13 patients by the parenteral administration of BPO-FLYS 100-400 mg./day. In all patients, skin tests with BPO-FLYS at the beginning of therapy were negative, but in three cases, slightly positive skin reactions were observed after eight to 31 days of therapy.

"Those patients who were undoubtedly hypersensitive to penicillin, and who had freshly experienced clinical allergic reactions, were capable of pursuing penicillin therapy under the protection of the haptene," said Dr. de Weck.

The findings will have to be confirmed by further clinical trials, now being conducted in research centers in Switzerland, France, and West Germany, he told MEDICAL TRIBUNE. "Obviously, we are not ready yet to put this into the hands of general practitioners, because there are still some problems. But the research work is going very well."

Dr. de Weck commented that the work offers a new approach to control of the immunologic system by depression of the formation of specific antibody.

"If it is feasible to depress the formation of antibody without impairing cell-mediated immunity, then we could have new possibilities in cancer therapy," he observed.

"We have to be able to identify the tumor antigen, and in some cases this knowledge is already available."

Coauthors were Drs. C. H. Schneider, H. Spangler, O. Toffler, and S. Lazary, all of the University of Berne.

Dr. David C. Marsh, an immunologist from Johns Hopkins University, working

at the Good Samaritan Hospital, Baltimore, said that his team's most recent work helped confirm the belief that allergies are genetically determined.

In exceedingly allergic patients, Dr. Marsh's group was able to demonstrate n

highly significant correlation between sensitivity to the ragweed allergen Ra5 and histocompatibility antigens of the cross-reacting group (HL-A7).

Coauthors were Drs. Wilma B. Blas, Susan H. Hsu, and Lawrence Goodfriend.

## WHO Experts List 6 Major Hazards To Health Found in the Environment

Medical Tribune World Service

GENEVA, SWITZERLAND—There are six major health hazards in the environment, according to World Health Organization experts meeting here. These are:

- Oxides of nitrogen, because of the unclear public health implications of these compounds in the ambient atmosphere.

- Polychlorinated biphenyls, because of their demonstrated toxicity and wide dissemination in water and packaging material.

- Asbestos, because of its demonstrated cancer-producing properties and widespread use for industrial, structural, and other commercial purposes.

An international program designed to develop environmental health criteria for the protection of man from this complex of environmental hazards was agreed upon at the meeting, which was under the chairmanship of Prof. Lars Friberg, of the Karolinska Institute, Sweden.

Called the father of inorganic chemistry, Baron Jöns Jakob Berzelius (1779-1848) was born in Sweden. Receiving degrees in both chemistry and medicine from the University of Uppsala, he taught pharmacy, medicine, and chemistry in Stockholm.

He discovered the elements selenium, thorium, and cerium and isolated biliverdin, a green pigment formed from bilirubin by oxidation. He introduced the present system of writing chemical symbols and formulas.

Sweden issued the stamp in 1939 to honor the 200th anniversary of the Royal Academy of Science. This year marks the 125th anniversary of Berzelius' death.

Text: Dr. Joseph Klar

Stamp: Mihluk Publications, Inc., New York

## A Microbicide Douche

Clinically Effective  
Trichomoniasis  
Nitrofurantoin

## BETADINE DOUCHE

for trichomoniasis

infectious vaginitis

nitrofurantoin

infectious vaginitis

## Experts Hail New FDA Food-Labeling Policy

Continued from page 1  
helpful to both doctor and patient by providing information that was formerly unavailable.

"Knowing about the fat content of food, whether it is polyunsaturated or not, and the calorie content will enable many patients to follow their physicians' advice more carefully," Dr. Calloway observed.

Nutritionists agreed that in managing malnutrition, cardiovascular disease, obesity, and sodium intake, the new labeling practices should close the gap between the physician's biochemical training and the existing lack of information on the food-stuffs his patients may be choosing.

"Malnutrition, the question of sodium intake, and the galloping mortality rate from cardiovascular disease," said Dr. Mayer, "are among the problems that require a definite stand on nutrition by physicians. In many cases, patients need to discuss with their physicians exactly what they have been eating, and many patients have not been doing this. This relabeling is going to make it much easier for patients to follow recommendations, and much easier for physicians to be direct about what the patient should or should not eat."

He emphasized that the new labeling practices will make it easier for physicians to learn more about nutrition.

"Now there will be no reason not to know, for example, which fats are polyunsaturated and which are not," he said. "The physician can go into his own kitchen from now on and find out for himself."

Dr. Mayer continued: "In a study which I recently did in the Boston area concerning the level of physician information on nutrition, I found out that most doctors remember their biochemistry quite well, but when it comes to applying their knowledge in a practical way to food, there is a decided gap. I think this relabeling plan will emphasize the need for more information."

### Jackson, With History Of Violence, Is Calmed By Amphetamine Therapy

Continued from page 1  
peared within an hour, Dr. Corson reported. With continued amphetamine therapy, Jackson welcomed the approach of laboratory personnel, even whimpering for further petting. He became nonaggressive with dogs previously attacked, and showed rapid learning in the pavlovian conditioning situation.

After six weeks of drug-facilitated psychosocial therapy, medication was withdrawn. Although the hyperkinesis reappeared, there was no recurrence of violent behavior and the dog did not forget what had been learned in the conditioning experiments.

"This has interesting implications for the learning of hyperkinetic or violent children in school under the influence of stimulants," Dr. Corson said. "Insofar as it is valid to extrapolate from animals to humans, this suggests that what such children learn in school while medicated with amphetamine they would tend to retain later."

#### Low Hyperkinesis Persisted

An additional two months of amphetamine and psychosocial therapy for the dog brought a reduction in the hyperkinesis that persisted even after withdrawal of the drug.

Dr. Corson found that dosages required for control of violent behavior were the same for dextroamphetamine or the levo isomer. By contrast, the control of hyperkinesis required four times as much levamphetamine as dextroamphetamine.

The differential effects of the two isomers, he commented, "would suggest the involvement of a dopaminergic system in violent behavior and primarily a noradrenergic system in hyperkinesis."

The investigator noted that genetic factors may have played a role in the behavior of this dog. All of its five littersmates exhibited similar behavior patterns.

tion and discussion, with benefits to both patients and doctors."

David Call, Ph.D., Professor of Food Economics, Graduate School of Nutrition, Cornell University, and a member of the FDA Commissioner's Food Advisory Committee, agreed that the new regulations may permit more specific advice from physicians and prompt more questions from patients. He noted, however, that they also may eliminate many questions now brought to physicians.

#### Won't Be Asking Doctors

"The specific prohibitions about saying certain things about food, if they are implemented, should clear up a lot of questions in consumers' minds so they won't be asking their doctors. They won't have to come to their doctor and say, 'Is it true that this food will cure cancer (or heart disease or something else)?" because that kind of misinformation will no longer be permitted," he said.

In addition to consumer-oriented information, such as serving size and servings per container, whenever a nutritional claim is made for a product the new regulations will require notice of calorie, protein, carbohydrate, and fat content as well

as percentage of U.S. Recommended Daily Allowances (RDA) of protein, vitamins, and minerals.

The RDA replaces the Minimum Daily Requirements as the official measurements of nutritional intake. Generally, they nearly double the standards of vitamins A, B<sub>1</sub>, B<sub>2</sub>, niacin, B<sub>6</sub>, folacin, pantothenic acid, B<sub>12</sub>, biotin, C, D, E, and K, and calcium, chlorine, iron, magnesium, phosphorus, potassium, sodium, sulfur, copper, fluorine, iodine, manganese, and zinc.

The combination of final regulations, tentative orders, and proposals put forth by the FDA will require listing of percentages of vitamins A, C, thiamin, riboflavin, and niacin as well as calcium and iron.

Manufacturers will also be allowed, but not required, to indicate the food's content of cholesterol, sodium, and polyunsaturated, saturated, and other fatty acids.

According to an official statement, "in taking this action the FDA is not taking a position on the scientific debate surrounding the role of fat consumption in heart disease. Consumers, however, should be able to identify foods for inclusion in physician-recommended fat-modified diets."

The new regulations would define as a

dietary supplement any item containing 50-150 per cent of the U.S. RDA of vitamins and minerals, require disclosure of their contents, and prohibit claims that they can prevent, cure, or treat disease. Any product exceeding 150 per cent of the RDA must be labeled and marketed as a drug.

#### Based on '68-70 Hearings

The U.S. RDA and supplemental dietary regulations are based on the Special Dietary Food Hearings conducted by the FDA during 1968-70.

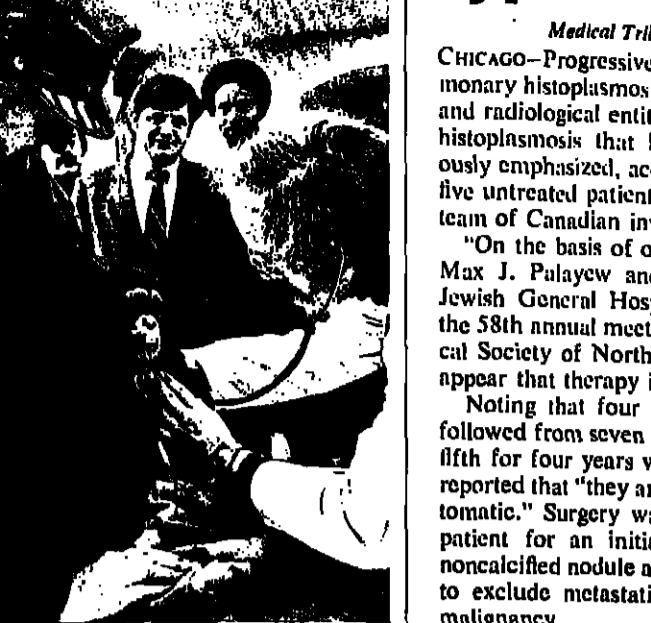
According to the head of the FDA, Dr. Charles C. Edwards, the regulations will implement virtually all the labeling recommendations of the White House Conference on Food, Nutrition, and Health.

He stressed that professionals must help consumers "understand and utilize the new labeling information."

"As the program gets under way," he said, "labels will begin routinely bearing information never before seen by the average consumer. It is important for all of us to make every effort to inform consumers on how to use this new labeling to the benefit of themselves and their families."

All of the FDA's actions are scheduled to be finalized within six months of their appearance in the *Federal Register* on January 19.

## Care for Zoo Animals



Pathologists from New York Medical College, currently conducting comparative studies of diseases shared by man and beast, will provide medical care for N.Y.C. zoo animals that are ill or injured. Above, Edward Garner, D.V.M., examines chimp at Central Park Zoo.

## Type of Histoplasmosis Needs No Treatment

### Medical Tribune Report

CHICAGO—Progressive, multinodular pulmonary histoplasmosis is a distinct clinical and radiological entity in the spectrum of histoplasmosis that has not been previously emphasized, according to a study of five untreated patients reported here by a team of Canadian investigators.

"On the basis of our experience," Drs. Max J. Palayew and Harold Frank, of Jewish General Hospital, Montreal, told the 58th annual meeting of the Radiological Society of North America, "it would appear that therapy is unwarranted."

Noting that four of the patients were followed from seven to nine years and the fifth for four years without therapy, they reported that "they are all well and asymptomatic." Surgery was performed in one patient for an initial solitary enlarging noncalcified nodule and in another patient to exclude metastatic spread of thyroid malignancy.

The investigators commented that their experience "would tend to support a most conservative approach to the patient with multiple histoplasmoses even in the face of growth and/or cavitation."

They emphasized that lack of awareness of this entity can lead to the needless

risks of thoracotomy and/or amphotericin B therapy.

Discussing the radiologic findings, the physicians said that, regardless of the initial radiologic presentation, all five patients subsequently developed multinodular parenchymal changes; two developed typical central calcification, and two showed cavitation at varying stages of their evolution. Nodules also showed both increase and decrease in size during follow-up examination. Some disappeared while others were developing.

"This variable radiologic picture remains somewhat puzzling in terms of

## Doppler Ultrasound Valuable In Detecting Venous Occlusion

### Medical Tribune Report

PHILADELPHIA—Doppler ultrasound is an "excellent" device for the detection of lower limb venous occlusion and is comparable in effectiveness with iodine-125 fibrinogen test, a relatively new diagnostic procedure, investigators from the Oklahoma City VA Hospital reported here.

A study of 52 patients demonstrated that there is no statistical difference between the two procedures in sensitivity in the detection of deep venous thrombosis, the investigators told the 17th annual meeting of the American Institute of Ultrasound in Medicine. Doppler ultrasound, however, has the advantage of being atraumatic, rapidly done, and not subject to interference from previous isotopic procedures, said Harold Poehlmann and Drs. Ross E. Brown and James M. Hartack.

In the I-125 fibrinogen test, they explained, the agent is administered intravenously and after a two-hour interval uptake counts are made with a scintillation counter. The probe is placed over a minimum of seven marked points following the deep venous drainage in the leg. An abnormal test is determined by a 20 per cent increase in counts at two consecutive points on the same leg on the same day and sustained for two days.

In the Doppler technique, they said, spontaneous venous flow is detected when the transducer is placed over a vein in the leg and this results in a cyclic blowing sound regulated by the respiratory cycle.

#### Flow Velocity Increased

"When a group of muscles are contracted or are squeezed, an additional quantity of blood flows into the venous system, momentarily increasing the velocity of venous flow," they remarked. "The sound derived from the increased velocity is called augmented flow. This is present when the deep venous system is patent."

"When the system is occluded, there is no augmented flow, an indication of venous obstructive disease."

The 52 patients were seen on seven consecutive days with a total of 728 limbs examined, they reported.

Three limbs were found with no augmented venous sound and abnormal I-125 tests were also noted. Venography confirmed these results.

Four limbs were found where the augmented venous sounds were not as loud as would normally be expected from a totally unobstructed vein. Of these four limbs, two were confirmed by I-125 as being equivocal, it was reported. The two other limbs were not followed by I-125 because of interference from a liver scan given during the same period. No venography was performed, and the patients completed an uneventful postoperative recovery.

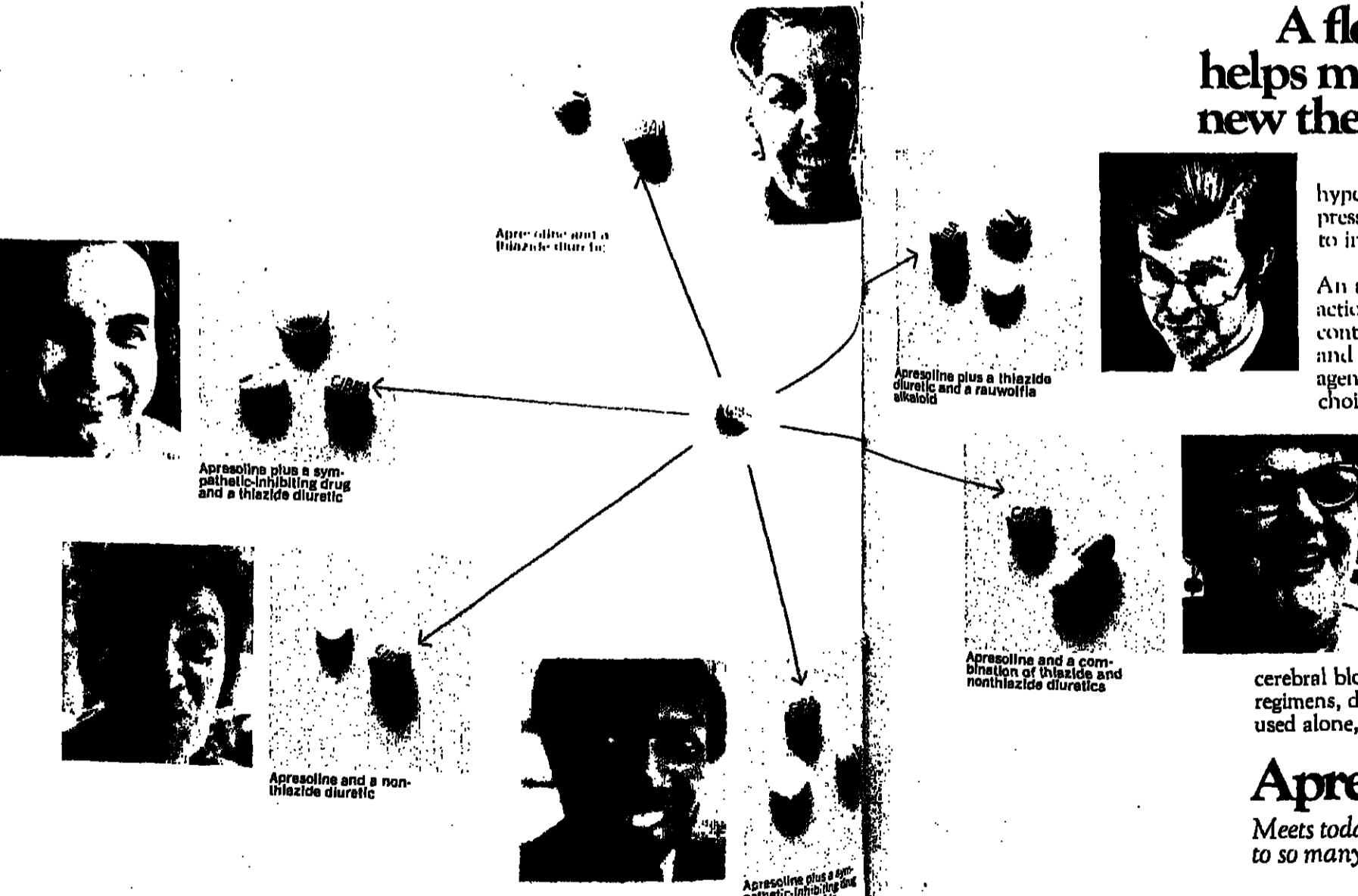
Of the remaining limbs, the investigators reported, 721 were normal on Doppler examination but two yielded equivocal results with I-125; these results, however, returned to within normal limits by the end of seven days.

An apparent disadvantage of Doppler ultrasound, the investigators noted, "is that all of the deep venous system on the calf must be occluded before the augmented flow is lost or the popliteal must be involved."

The three main veins in the calf, they pointed out, are the anterior tibial, posterior tibial, and popliteal. "If a single vein is occluded, the augmented sound may be heard, although it may be diminished. Once the thrombosis has progressed from a single vein into the popliteal, the augmented sound will be lost."

## Apresoline...an antihypertensive idea whose time has come

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Early and more vigorous treatment of hypertension. More adequate control of blood pressure. Antihypertensive regimens closely molded to individual requirements.

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Apresoline differs from other available antihypertensives in that it appears to act directly on the arterioles where diastolic blood pressure is ultimately controlled. By relaxing arteriolar smooth muscle, it decreases peripheral vascular resistance—decreases arterial pressure.

Apresoline also helps increase renal blood flow and maintain glomerular filtration, and to maintain or increase cerebral blood flow. When Apresoline is added to existing regimens, dosages of each drug are usually lower than when used alone, thus tending to reduce risk of side effects.

### Apresoline (hydralazine)

Meets today's needs because it can contribute so much to so many antihypertensive regimens

#### Apresoline<sup>®</sup> hydrochloride (hydralazine hydrochloride)

TABLETS

INDICATIONS

CONTRAINDICATIONS

WARNINGS

PRECAUTIONS

ADVERSE ACTIONS

OVERDOSE

reactions are reversible upon withdrawal of therapy, but long-term treatment with steroids may be necessary. An L.E. cell preparation is indicated in the presence of any unexplained symptoms.

Use MAO inhibitors with caution.

Although there has been no adverse experience with Apresoline in pregnancy, this drug should be used only when, in the judgment of the physician, it is deemed essential to the welfare of the patient.

Use cautiously in suspected coronary artery disease, peripheral vascular disease, and valvular rheumatic heart disease.

WARNINGS

May produce anorexia, headache, and dizziness.

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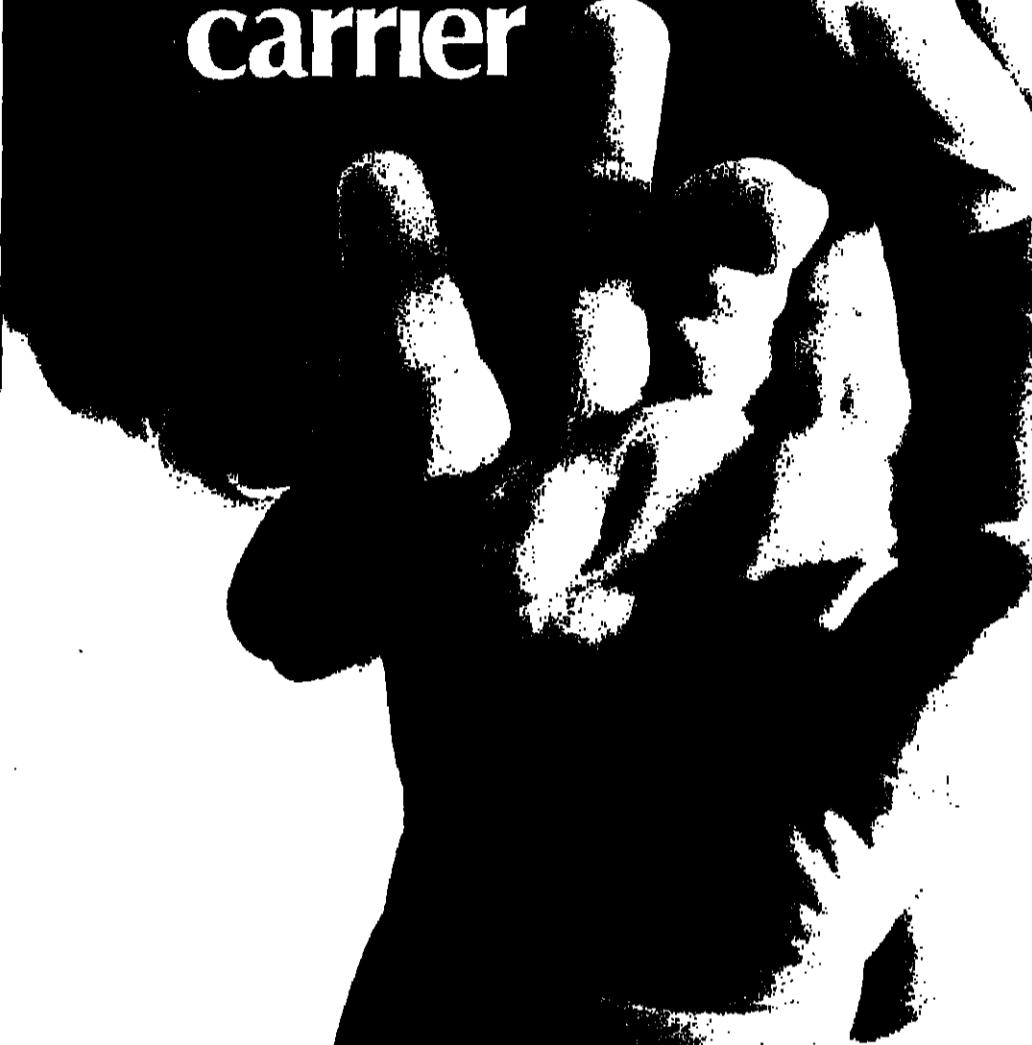
Use cautiously in suspected coronary artery disease, peripheral vascular disease, and valvular rheumatic heart disease.

WARNINGS

May produce anorexia, headache, and dizziness.

PRECAUTIONS

# the caring hand is not a carrier



The nurse's hand washed with phisoHex® is an important part of the anti-Staph protection for the newborn. The protection can be maintained throughout the infant's stay in the hospital nursery by having nurses wash their hands with phisoHex before and after handling each infant.

The physician can maintain this antibacterial protection at home by prescribing the use of phisoHex for mother's hands before handling the baby. phisoHex creates a bacteriostatic film on skin. There it remains to inhibit growth of microorganisms.

And nonalkaline, hypoallergenic phisoHex is kind to skin. Won't tend to dry or irritate, even when used frequently.

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to help take the Staph problem off your hands

**phisoHex®—Brief Summary**  
sudsing antibacterial soapless skin cleanser  
phisoHex contains a colloidal dispersion of hexachlorophene 3% in a stable emulsion consisting of emulsifier sodium octylphenoxethoxyethyl ether sulfonate 50%, petroleum 7%, lanolin cholesterol 0.7%, methylcellulose, polyethylene glycol, polyethylene glycol monostearate, lauryl myristyl diethanolamide, sodium benzoate, and water. pH 5.0 to 6.0 is adjusted with hydrochloric acid. All ingredients w/w.

**Actions:** phisoHex has bacteriostatic action against staphylococci and other gram-positive bacteria. Cumulative antibacterial action develops with repeated use.

**Indications:** phisoHex is indicated for use as a surgical scrub and a bacteriostatic skin cleanser. It may also be used for washing to control an outbreak of gram-positive infection in the nursery when good hospital practice has been inadequate as a total program of infection control. It should be used only as long as necessary for infection control.

**Contraindications:** phisoHex should not be used on burned or denuded skin. It should not be used as an occlusive dressing, wet pack, or lotion. It should not be used routinely for prophylactic total body bathing. It should not be used as a vaginal pack or tampon, or on any mucous membranes. phisoHex should not be used on persons with sensitivity to any of its components. It should not be used on persons who have demonstrated primary light sensitivity to halogenated phenol derivatives because of the possibility of cross-sensitivity to hexachlorophene.

**Warnings:** *Rinse thoroughly after use*, especially from sensitive areas such as the scrotum and perineum. If left in contact with burned or denuded skin or mucous membranes, sufficient hexachlorophene may be absorbed to cause toxic symptoms. Infants, especially premature infants or those with dermatoses, are particularly susceptible to hexachlorophene absorption. Systemic toxicity may be manifested by signs of stimulation (irritation) of the central nervous system, sometimes with convulsions. *phisoHex should be discontinued promptly if signs or symptoms of central instability occur.* Experimental and clinical evidence indicates that hexachlorophene toxicity is reversible.

In a small number of reported cases, fatal intoxications from hexachlorophene have occurred. These cases include misuse of 3% hexachlorophene on burned skin or exposure to a powder accidentally containing approximately 6.5% hexachlorophene. Examinations of brain tissue in some of these cases revealed vacuolization like that which can be produced in newborn experimental animals following repeated topical application of 3% hexachlorophene for 90 days.

phisoHex is intended for external use only. If swallowed, phisoHex is harmful especially to infants and children. phisoHex should not be poured into measuring cups, medicine bottles, or similar containers since it may be mistaken for baby formula or other medications.

**Precautions:** phisoHex suds that get into the eyes accidentally during washing should be rinsed out promptly and thoroughly.

**Adverse Reactions:** Dermatitis and photosensitivity. Sensitivity to hexachlorophene is rare; however, persons who have developed photoallergy to similar compounds also may become sensitive to hexachlorophene.

In persons with highly sensitive skin, the use of phisoHex may at times produce a reaction characterized by redness and/or mild scaling or dryness, especially when it is combined with such mechanical factors as excessive rubbing or exposure to heat or cold.

**Treatment of Accidental Ingestion:** The accidental ingestion of phisoHex in amounts from 1 to 4 oz. has caused anorexia, vomiting, abdominal cramps, diarrhea, convulsions, hypotension and shock, and in several reported instances, fatalities. (See Prescribing Information for detailed treatment.)

**How Supplied:** phisoHex is available in unbreakable plastic squeeze bottles of 5 ounces, 1 pint, and in plastic bottles of 1 gallon.

For detailed DIRECTIONS, consult Prescribing Information.

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# Medical Tribune

and Medical News

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## Bravo, Commissioner!

We have on previous occasions congratulated Dr. Charles C. Edwards, Commissioner of Food and Drugs, and we do so again, now that the FDA is introducing the changes in good labeling practices that will permit consumers to know the content of processed foods (see page 1). On this occasion we can clearly show the difference in the position taken by the FDA under Dr. Edwards' leadership from that assumed six years ago under another commissioner. We reprint the following editorial that was published in MEDICAL TRIBUNE, May 22, 1967.

## The FDA and Fats in the Diet

**T**HE PROFESSOR OF NUTRITION at the Harvard School of Public Health, Jean Mayer, Ph.D., D.Sc., sharply revived the question of labeling edible fats, oils, and fatty foods to show percentages of unsaturated and saturated fatty acids. In his address before the Division of Environmental Sciences of the New York Academy of Sciences, Dr. Mayer went further than that. He said, "It is unfortunate that our Federal Government, which has already dragged its feet to a scandalous extent as regards action against cigarette smoking, is equally negligent as regards saturated fat, with the Food and Drug Administration and its new director refusing to allow advertising claims which would emphasize the cardiovascular advantages of polyunsaturated fatty acids and, therefore, encourage industrial concerns to change their processing customs to encourage a change in the nature of the fats used."

For many years an association has been noted between the incidence of coronary artery disease and the levels of blood cholesterol and other lipids, and these in turn have been related to the dietary intake of particular fats. In 1961 the American Heart Association called for "reasonable substitution" of polyunsaturated for saturated fats, because this would help reduce blood cholesterol levels and because the incidence of coronary artery disease in our country is unreasonably high. In 1965, the A.H.A. made stronger recommendations. It noted that "in most persons, but not all, the level of cholesterol and other fats in

the blood can be decreased and maintained at a lower value by conscientious and long-term adherence to a suitable diet." The A.H.A. urged for most people a significantly decreased intake of saturated fat and a significantly increased intake of polyunsaturated fat, with polyunsaturated fats being substituted for saturated fats in the diet wherever possible.

But in 1959 the FDA ruled that labeling of a food that implied that consumption of polyunsaturated fats could prevent or treat heart or artery disease was a misdemeanor. In 1965 the FDA invited interested parties to file statements on a proposed regulation that a food represented as of special dietary use in the intake of fatty acids bear a label listing accurately the number of grams of saturated, monounsaturated, and polyunsaturated fatty acids contained in an ordinary serving and in 100 gm. This was done at the request of the American Diabetes Association and six prominent clinicians in heart disease and nutrition.

Early in 1966 Dr. James L. Goddard, Food and Drug Commissioner, rejected the proposal and stated that it was the agency's position that manipulation of blood cholesterol levels through diet is not "conclusively accepted by scientists as the best way to prevent, treat, or control heart or artery disease." It is this ruling of Commissioner Goddard that Dr. Mayer objects to. We object to it, too, and find it disturbing that the FDA has in its power to make and enforce such a decision in the face of contrary opinion based on abundant research by expert investigators and physicians.

## Cigarettes and Women

**A**CCORDING TO A STUDY by Dr. David M. Spain and his colleagues, 62 per cent of women dying from coronary heart disease were heavy cigarette smokers; this was true of only 28 per cent of women dying from other causes. The incidence of lung cancer among women has also risen with an increase in their smoking habits.

And now the latest annual report to Congress by the Public Health Service on the consequences of smoking emphasizes that risk to the fetus falls in that area, but we call upon women to discontinue smoking not for that reason alone.

## The Hyperkinetic Dog

**E**XPERIMENTAL QUOTE: "We do not wish to leave the impression that all violent behavior can be eliminated with the help of drug therapy. Psychosocial therapy should be tried before any drug administration is instituted. Our studies suggest that in some types of violent behavior which cannot be controlled by any



"Dr. Parker, Internist; Dr. Walski, nephritis specialist; and Mr. Forsham, our expert on insurance forms."

ing. Is it so painful to present the truth? Does Dr. Edwards believe that his organization's "expertise" with respect to the utilization of drugs would be banned as evidence in any court in this country? I think not.

The indirect threat of economic sanction by litigation is a technique well-known in government circles, and with inflamed consumers and their legal counsel ready to take up the cause at a moment's notice, the FDA need take no action other than "regulating the drug." The credibility gap persists—only the camouflage wears thin.

C. EARL HILL, M.D.  
University of Maryland

## The Vas Rejoined

Editor, MEDICAL TRIBUNE:

In a recent issue you reported that vasectomy "has become an increasingly popular and widely accepted means of birth control in the United States." This is indeed true. Vasectomies increased from about 50,000 yearly during the 1960s to some 750,000 in 1970.

Your article then rather deplored the number of people who thought that vasectomies were reversible and suggested that "medical and allied professions make certain that persons seeking a vasectomy fully understand the permanency of the operation."

No man or woman considering a sterilization should assume that it could be easily reversed, and must think of it in terms of being permanent. However, it should also be pointed out that, depending on the techniques used for the sterilization and on the very special skills of the surgeon performing the reversal and also, perhaps, on luck, it is possible to restore fertility by rejoining the severed tubes or vas. It would not be fair, therefore, to disapprove of sterilization on the grounds that it is totally irreversible.

In my book on sterilization I quote Donald A. Goodwin, M.D., head of urology at the U.C.L.A. Medical Center, as saying that in the hands of experienced and well-trained urologists one should expect to achieve up to 90 per cent success in restoring fertility following vasectomy. Dr. John W. Dorsey of Long Beach reported a success of over 80 per cent in a series of over 100 cases and Elmer Bell of Los Angeles has reported 85 per cent success in rejoining the vas so that sperm cells once again appeared in the semen.

H. CURTIS WOOD, JR., M.D.  
Fort Washington, Pa.

## FDA—Drug Regulation

Editor, MEDICAL TRIBUNE:

I was amused to read Dr. Charles C. Edwards' response to the question whether his administration was regulating drugs or doctors [interview, MEDICAL TRIBUNE, January 10].

In attempting to examine his response logically, we must reason that even the FDA cannot regulate drug efficacy, mode of action, chemical composition, side effects, etc. The FDA can and does regulate its manufacture, purity, and distribution.

When you regulate its use, you do facto regulate the individual who effects its ultimate distribution to the consumer—the prescribing physician. A rose remains a rose despite Dr. Edwards' hedge.

**Editor's Note:** The Supreme Court decision on abortion has dimmed the significance of controversy. Correspondence on the subject must therefore now be closed.

## Gut Flora Thought to Hold Key To Diet-Colon Cancer Relation

Medical Tribune Report

ATLANTA, GA.—A theory that relates cancer of the colon to diet—with the gut bacterial flora serving as a "vital intermediary" in the relationship—was outlined by a British investigator here at an International Conference on Anaerobic Bacteria.

Dr. M. J. Hill, of the Wright-Fleming Institute, St. Mary's Hospital Medical School, London, said the search for a dietary factor in colon cancer has been under way since 1967, when epidemiologic studies showed a much lower incidence of this malignancy in Japan, East Africa, and India than in Western Europe or North America.

Various research groups, he added, have suggested that such differences in incidence might derive from different intakes of food elements ranging from fat and protein to refined carbohydrate and fiber.

"Our studies, based on World Health Organization statistics, show the incidence of colon cancer to be strongly correlated with the amount of dietary fat and animal protein and not at all with dietary fiber," Dr. Hill told the conference, which was sponsored by the Center for Disease Control, the Upjohn Company, and Emory University.

### Correlation Coefficients Listed

The correlation coefficient between bound fat and incidence of colon cancer cited by Dr. Hill was a high 0.88; a strong correlation was also found between bound fat and breast cancer (correlation coefficient 0.80). The correlation coefficient between animal protein and incidence of colon cancer was 0.87 (0.79 for breast cancer).

By contrast, dietary fiber appeared to have little or no correlation with either form of cancer, and refined sugar showed coefficients of only 0.32 and 0.50.

Noting that the previous hunt for preformed carcinogens in the diet had not produced any adequate explanation for the diet-colon cancer correlation, Dr. Hill said he and coinvestigators began with the hypothesis that the gut bacteria might play a role as intermediaries. They postulated that:

- Cancer of the colon is caused by production of carcinogens and/or carcinogens by gut bacteria from dietary components or from intestinal secretions produced in response to the diet.
- The nature of the diet affects the composition of the intestinal bacterial flora and determines the substrates available for bacterial metabolism.
- Since the diet controls the intestinal

flora, the substrates available for carcinogen production, and also the physiologic conditions within the gut, this would explain the correlation between diet and the incidence of colon cancer.

Fat was chosen as the dietary component most likely to be concerned, Dr. Hill pointed out, because the amount of dietary fat determines the concentration of steroids in feces "and many acid steroids have been claimed to be carcinogenic."

The team's working hypothesis was that the amount of dietary fat determines both the concentration of bile acids and cholesterol in the large intestine and the bacterial flora acting on these steroids and that bacteria can produce carcinogens and/or carcinogens from the biliary steroids.

Fecal specimens from people living in areas of high and low incidence of colon cancer were then examined for bacterial flora and steroid content.

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When the two types of specimens were compared, the investigators found that feces from people in low-incidence areas had fewer anaerobic gram-negative *Bacteroides* organisms and more enterococci than did feces from people in high-incidence areas. Also, specimens from the low-incidence areas had a much smaller amount of fecal steroid (both acid and neutral) and such fecal steroids were much less bacterially degraded.

"Considering these results in the light of our working hypothesis," Dr. Hill said, "the amount of presumed substrate available for carcinogen production was greater in the high-risk groups and the degree of bacterial action was also greater."

Chemical studies have yielded support for the theory that bacteria can produce a carcinogen from biliary steroids and possibly from amino acids, according to Dr. Hill.

One area of investigation has been the bile acids synthesized in the liver—cholic acid and chenodeoxycholic acid. Bacterial dehydroxylation of cholic acid produces deoxycholic acid, a substance considered carcinogenic by some scientists.

Although its apparent carcinogenicity in rats has been disputed, Dr. Hill commented that "there is an extremely good correlation between the mean fecal concentration of deoxycholic acid and the incidence of colon cancer" in the fecal specimens examined from low-incidence and high-incidence areas.

The possibility that bacteria might produce a polycyclic aromatic compound from the biliary steroids was also investigated by Dr. Hill's team. Four types of re-



On Growth Hormone

## THE MEDICAL TRIBUNE BULLETIN

### New Use for Eosinophils

MONTREAL—Dr. Thomas Hubscher, of Montreal Children's Hospital, reported that eosinophils were found to contain a soluble factor capable of inhibiting allergic histamine release from sensitized target cells—i.e., basophils and/or mast cells.

"And man is bountifully supplied with eosinophils," he commented. "The implication is that if we can isolate this substance in pure form and synthesize it, it could be a very productive drug with minimal side effects."

He spoke at an international conference on control mechanisms in reagin-mediated hypersensitivity, held in honor of Dr. Brum Rose, retiring allergist-in-chief of Royal Victoria Hospital and Professor of Experimental Medicine at McGill University.

Dr. Hubscher's coauthor was Dr. A. H. Eisen.

### Hyperlipoproteinemia

WIESBADEN, WEST GERMANY—The likelihood of hyperlipoproteinemia in parents can be forecast from a determination of total cholesterol and beta-cholesterol levels in newborn infants, according to a German investigator.

In addition, the cholesterol levels can indicate whether the child is likely to develop the disease in later life, said Dr. Horst Wengeler, of the Heidelberg University Hospital Department of Medicine.

Total cholesterol is determined in whole serum. After ultracentrifugation, the cholesterol level is determined in the low-density plus high-density lipoprotein fraction. From this determination, the cholesterol level present in the high-density lipoprotein fraction is subtracted. This yields the beta-cholesterol level.

The disease was diagnosed in 13 of the parents of over 150 newborns in whose umbilical cord blood high total cholesterol and beta-cholesterol levels had earlier been detected, he told a meeting of the German Society for Internal Medicine.

His co-workers were Drs. Heiner Greten and Mathias Wagner.

### Drug for Sex Offenders

SAN REMO, ITALY—The libido-dampening effect of cyproterone acetate is having an impact on judicial decisions in Germany and Switzerland, Dr. E. Rainer, of the Medical Division of Schering S.p.A., Milan, told MEDICAL TRIBUNE at an International Congress on Sexology.

In Switzerland, reduced sentences have been imposed in some sex offense cases when the offender agreed to undertake treatment with the drug.

"In Germany, where the sexual delinquent can get his freedom by allowing himself to be castrated, treatment with cyproterone acetate has been accepted by the Government as an analogue to the effects of castration," said Dr. Rainer.

Dr. P. Saba reported to the congress that the drug proved successful in the treatment of eight oligophrenic, cerebropathic patients suffering from hypersexuality characterized by exhibitionism, aggressivity, and continuous masturbation, at the Psychiatric Hospital of Volterra, Italy, where he is chief physician.

### A Suit Over Drugs

OSAKA, JAPAN—Fifty-three victims of subacute myelooptic neuropathy have filed suit here for \$4,800,000 in damages from the Japanese Government and seven pharmaceutical companies that imported, produced, or sold drugs containing lodo-chlorhydrquin, the suspected cause of their disease.

Counsel for the plaintiffs said that the suit is intended to clarify the responsibility of the Government for allowing the companies to sell the drug without confirming its safety.

The team also sets up and measures

## Team Reduces Cord Patient Hospital Stay

Medical Tribune Report

DOWNEY, CALIF.—The coordinated action of a team of several professionals and paraprofessionals in the treatment of patients with severe spinal cord injuries has drastically reduced their length of stay in the hospital, according to Dr. Frederick N. Elliott, assistant medical director of Rancho Los Amigos Hospital here.

If such patients are admitted within two weeks of their injury, the average length of stay is 100 days less than for those patients who are admitted after that time, after having been treated elsewhere. In terms of cost, this means a saving of some \$20,000, he said.

Dr. Elliott reported that the entire team assigned to a particular patient—including student nurses, technicians, medical students, and nurses aides as well as the physician, nurses, psychologist, social worker, or physical therapist—join together in a conference on diagnosis and on frequent subsequent conferences on treatment progress and then discharge planning.

Special emphasis is placed on shifting his position frequently and prevention of decubitus ulcers.

The team also sets up and measures

goals, which also include prevention of loneliness as well as good physical care. "In this era we expect more than mechanical treatment of their disease," said Dr. Elliott. "We're both saving money and treating the patients much better."

Increased patient satisfaction has also gone along with increased satisfaction among the professionals and paraprofessionals, resulting in a 50 per cent reduction in turnover in team members. An added advantage has been the educational advantage to students in being so closely connected with a team.

### Philippine Dogs Vaccinated In Effort to Deter Rabies

Medical Tribune World Service

MANILA—House-to-house teams have vaccinated an estimated 80 per cent of the Philippines' canine population in a country-wide campaign to stamp out rabies.

A recent study showed that an average of 250 Filipinos contract rabies each year but that from 100,000 to 150,000 persons annually require preventive vaccinations after being bitten by suspect animals.

Because of the comparatively large group, the patient can also be taken care of around the clock.

Special emphasis is also placed on shifting his position frequently and prevention of decubitus ulcers.

The team also sets up and measures

Medical Tribune

# HYPERTENSION BULLETIN

ACIBA SERVICE

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EXACTING, UNFAMILIAR TASKS, in which failure may mean punishment, can induce arterial hypertension in the squirrel monkey. Another set of tasks can reverse the condition in the same animal. But not all monkeys subjected to precisely the same conditions develop hypertension. Thus for them, as well as for some human beings—strong emotional effects may induce organic disease.

This report, a preliminary one from the new Specialized Center of Research in Hypertension at Harvard Medical School, casts new light on a theory first proposed by the Harvard physiologist Walter B. Cannon, who published his classic text, *Bodily Changes in Pain, Hunger, Fear, and Rage*, in 1929.

Harvard researchers are nearing their 18th month of work on a long-term collaborative study to document physiologic mechanisms that promote organic disease in subhuman primates. One physiologist gives a capsule summary of their relationship to Cannon's theories: "Cannon had a strong interest, and produced some striking leads, in various areas of psychosomatic medicine. But he had few experi-

### ORIGINS OF HYPERTENSION: you're driving me nuts...

mental data. Our work so far finds no instance where he was completely wrong in his assumption. But our work also shows that many uncertainties remain."

The long-range goal of this study, supported by a grant from the National Heart and Lung Institute, is to find means to prevent and treat human hypertension.

Dr. A. Clifford Barger, the Hypertension Center's general director and principal investigator, points out that about 30,000 people in this country have some

form of cardiovascular disability and the Public Health Service figures that perhaps \$30.5 billion is spent annually in patient care for this disability. This includes direct costs of 5.1 billion dollars, and indirect costs of 25.4 billion dollars.

"If we were able," said Dr. Barger, "to postpone the onset of cardiovascular disease for five to 10 years—not an unreasonable goal for the next decade, provided our research momentum is maintained—the savings would be many billions of dollars."

At present, investigators of human hypertension are confronted by a complex set of unknowns, according to Dr. J. Alan Herd, Associate Professor of Physiology. To clear the way, the Harvard group is attempting to document the role of the environment—and indirectly, the emotions—in producing blood pressure elevation in laboratory monkeys.

"We chose squirrel monkeys because we needed totally naive subjects on whom we could impose a set of completely controlled and unfamiliar circumstances. Our monkeys sit alone in a chair in a very small chamber, responding to flashes of

continued on page 24

THE NATION IS TRYING to get an effective hypertension detection and treatment program under way, ultimately to cut down the massive social costs of cardiovascular disease; but there is a stricture in the channels of control; many people do not follow back for follow-up examinations. Why not?

Dr. Frank A. Finnerty, chief of cardiovascular research at Georgetown University Medical Division, District of Columbia General Hospital, put the question to himself when he found that many people were dropping out of his inner-city hypertension clinics. He organized a study to find out why—and the upshot has been a tactical and structural reorganization of clinical facilities.

In 1970 the Veterans Administration Cooperative Study Group on Antihypertensive Agents found that control of blood pressure in patients with diastolic pressures ranging from 90 to 114 mm. Hg significantly lowered morbidity and mortality. The NHLI then decided to set up a cooperative, nationwide study to discover whether these findings hold true for the population at large.

D. C. General moved into the study, prepared to use its several established clinics as examination centers for the Metropolitan Washington Regional Hypertension Detection and Follow-up Program.

"The incidence of hypertension among inner-city blacks," said Dr. Finnerty, "is high, approximately 40 per cent, compared with the 12-15 per cent in the general pop-

## RETURN OF THE CLINIC DROP-OUTS

ulation. It occurs earlier and is more severe. Among blacks, screening should start at age 25—and it isn't uncommon to find hypertension in teen-agers. Blacks seldom get coronary but often get strokes. No one understands why. Strokes are as common in women as they are in men, and in the D. C. population it is not unusual to see women in their early 30s who have had strokes. We don't know whether this is a racial difference or a result of the socioeconomic stress in this area.

"We learned quickly that we couldn't use standard epidemiological techniques for screening. In spite of support by community leaders in the census tracts and considerable favorable publicity in the local media, house-to-house screening turned out to be dangerous. On the first day of canvassing, a female member witnessed a rape. On the second day, some one tried to rape her."

So they set up screening centers in the largest supermarkets in each of three census tracts, and 61 per cent (6,480 of 10,560) of the residents of the tracts were screened in the markets. Nine hundred fifty-three were found to have pressures of 140/90 mm. Hg or higher, and these were invited to D. C. General for verification tests.

"We quickly learned that our first mistake was in scheduling appointments a week after the initial screening. About half

failed to show up. We were able to reduce this loss to 29 per cent by personal contact, and later to 5 per cent by making appointments within 24 to 48 hours. Each person who came to the clinic had two verification tests, and 296 were excluded because their diastolic pressures fell below 90 mm. Hg on the first or second visit. Along with dropouts, this left us with 284 patients for the study."

Dr. Finnerty and his colleagues supposed that the dropout rate was related to black suspicion of white professionals, to inadequate understanding of the seriousness of the disease, and to economic factors. But a good look exploded the assumptions.

"These patients had perfectly good reasons for not coming back to clinics. In the first place, each visit meant hours of waiting, an average of 2.5 hours before they were seen, and another 1.8 hours waiting at the pharmacy. This was on top of traveling time.

"When the patient did see the doctor, he got about seven and a half minutes of medical time. There was no real doctor-patient relationship. Not only was the doctor always in a hurry, but this is a teaching hospital and patients would see a different doctor at each visit, because of staff rotations.

"It's often assumed that clinic patients aren't motivated to get health care because they don't understand its importance. But the overwhelming percentage of the people in this study were perfectly aware that hypertension is a serious disease, and 56 per cent considered regular medical checkups important.

"We learned that the problem wasn't with the patients, but with how the patients were treated. After wasting a couple of days waiting around, patients say: 'The hell with it! Not even a bonus system would bring them back, and the next time we'd see them would be in the emergency room with a stroke or a coronary.'

Guided by the patients' complaints, procedures were changed. The Hypertension Clinic at D. C. General is kept open six days a week. Patients are seen by appointment. Every patient who is selected for follow-up is assigned to a physician and a paramedical health aide. At

every visit he sees the same physician and the same paramedical.

"If a patient misses an appointment, the health aide gets in touch to find out why. If it is a matter of a baby sitter or transportation, the aide finds a solution, even if it means that we arrange to pick the patient up and bring him to the clinic.

"For the most part, it's the paramedicals to whom the patients turn for information. They work under the supervision of nurses and use the doctors as consultants, but once a patient has been stabilized on medication, the aide follows the case, calling on the doctor only in the event of complications."

The Hypertension Clinic at D. C. General central clinic also offers comprehensive health care; the medical staff members act in the role of family doctors. The clinic phone is manned 24 hours a day, and there is a system for emergency services, outpatient care, and hospital admission.

"We bypassed the waiting time at the pharmacy by dispensing medication right in the clinic."

Once the clinic was operating for the benefit of the patients rather than for the convenience of the medical staff, comments Dr. Finnerty, the dropout rate fell from the high 42 per cent of 1966-1969 to 8 per cent.

In Dr. Finnerty's opinion, all clinics will have to be reorganized along these lines if "we are really going to treat and follow up patients with chronic diseases, such as hypertension." And he sees paramedicals as vital personnel in clinic staffs, contributing much more than the medical duties for which they are trained.

"Paramedicals will have to be brought into the system," he asserts. "They have to be legalized, have the right to third-party payments, and be covered by liability insurance. It's going to be difficult to persuade doctors that this concept isn't a threat to them. We can't force them to accept it. We can only show them, through repeated successful demonstrations, that paramedicals are the answer to overcrowded clinics and doctors' offices."

## reports from abroad

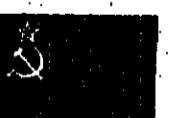
VARNA, BULGARIA—Electrosleep therapy combined with climatotherapy depresses blood lipid levels, according to a study by Prof. Dr. V. Sirkova, Director of Internal Medicine and Therapy, Institute of National Economy. Therapy depressed blood pressure, serum cholesterol, and beta-lipo-protein lipase activity in males and total lipid and triglyceride levels in females.



ULAN BATOR, MONGOLIA—A hypertension control program among various Mongolian nationals, aged 15 to 70 years, revealed: among 1,963 males, mean systolic blood pressure of 125.7, mean diastolic of 79.0; among 2,015 females, figures were 122.0 and 77.6, respectively. Diet for these peoples with common customs and traditions, is low in fruits and vegetables, high in sweets. Staple foods are meat—primarily fat mutton—and dried homemade milk products. Daily protein intake averages 109.5 Gm., 68-71 percent of which is of animal origin.



VARNA, BULGARIA—Patients with primary arterial hypertension as well as those with hypotension respond favorably to electrosleep therapy using low-frequency electric impulses, according to Prof. Dr. L. A. Studnizyna, of the Central Research Institute for Balneology and Physiotherapy, Moscow. Using this procedure, marked improvement was obtained in 96 per cent of 180 patients with hypotension and in 83 per cent of 135 with hypertension. Dr. Studnizyna reported at the third International Symposium for Electrosleep and Anesthesia.



MOSCOW—Study of arterial hypertension among 16,000 men aged 40-49 years revealed that arterial hypertension with increased systolic pressure only is not widespread: 0.7 per cent in the 40-44 year age group; 1.6 per cent in the 45-49 year group. Diastolic hypertension is more frequent: 10.1 per cent and 12.7 per cent, respectively, for the two age groups. Simultaneous rise in systolic and diastolic pressures occurred in 7.9 per cent of 40-44 year-olds and in 10.6 per cent of the older group.

# Two ways to treat moderate hypertension and why...



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Ser-Ap-Es should be used with caution in patients with advanced renal damage and cerebrovascular accidents. It should be discontinued at the first sign of mental depression.

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Esimil, an equally valuable yet different approach to moderate hypertension, makes sense for many patients because it anticipates future problems while helping to solve present ones.

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Postural hypotension may occur with the use of Esimil, particularly while the drug is being introduced. Like all antihypertensives, Esimil should be given with caution in the presence of severe coronary insufficiency or recent myocardial infarction.

early, effective control of hypertension can save lives

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Two ways to  
treat moderate  
hypertension

## Esimil®

guanethidine monosulfate 10 mg  
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## Ser-Ap-Es®

reserpine 0.1 mg  
hydralazine hydrochloride 25 mg  
hydrochlorothiazide 15 mg

**INDICATIONS**  
Ser-Ap-Es is recommended for all cases of hypertension, except the mildest and the most severe.

### CONTRAINDICATIONS

Known hypersensitivity, mental depression (especially with suicidal tendencies), active peptic ulcer, ulcerative colitis, digitalis intoxication, edema, pregnancy, and patients receiving electroconvulsive therapy, hydralazine. Hypertension in patients in general have a higher risk of intrapartum hypertension and other cardiovascular complications than normotensive patients. Reserpine-treated patients are not known to have a higher risk of hypertension than those with otherwise comparable hypertensive patients.

Preoperative withdrawal of reserpine does not assure that circulatory instability will occur. It is important to withdraw reserpine at least two weeks before surgery and consider this in the overall management, since hypertension has occurred in patients receiving rauwolfia derivatives. Antihypertensive agents (e.g., reserpine, guanethidine, etc., etc., norepinephrine) have been employed to treat severe hypovascular effects. Hydralazine stimulation produced by hydralazine can cause anginal attacks and ECG changes of myocardial ischemia. The drug has been implicated in the production of myocardial infarction. It must, therefore, be used with caution in patients with suspected coronary artery disease.

The "hyperdynamic" circulation caused by hydralazine may accentuate special cardiovascular hazards, particularly in patients with a history of increased pulmonary artery pressure in patients with mitral valvular disease. The drug may reduce the pressor responses to epinephrine. Postural hypotension may result from hydralazine but is less common than with reserpine. Drug-induced depression may persist for several months after drug withdrawal.

Hydralazine should be discontinued for at least two weeks before giving electroshock therapy. MAO inhibitors should be avoided or used with extreme caution.

Hydralazine administration of doses over 400 mg per day may produce in a few patients an arthritic-like syndrome leading to a clinical picture simulating acute systemic lupus erythematosus. A rare but serious reaction may occur at lower doses and symptoms and signs usually regress when the drug is discontinued, but long-term treatment with steroids may be necessary. Adverse reactions to hydralazine may be seen in patients with a history of hepatic coma. L.E. cells may be found in the blood of patients on the drug who are asymptomatic. An L.E. cell preparation is indicated if the patient has arthralgia, fever, testicular pain, continued malaise, or other unexplained symptoms.

Use MAO inhibitors with caution in patients receiving hydralazine. **Hydralazine**

There have been several reports, published and unpublished, concerning nonspecific small bowel lesions consisting of stenosis, with or without ulceration, associated with the administration of enteric-coated hydralazine. These lesions may occur with plain tablets or with coated tablets. These lesions may occur with enteric-coated potassium tablets alone or when they are used with nonenteric-coated thiazides for certain other oral diuretics.

These lesions have caused obstruction, hemorrhage, and perforation. Surgery was frequently required and deaths have occurred.

Available information tends to implicate enteric-coated hydralazine, without lesions of this type also occurring spontaneously. Therefore, coated potassium-containing formulations should be administered only when indicated and should be discontinued immediately if any evidence of nausea, vomiting, or gastrointestinal bleeding occurs.

Coated potassium tablets should be used only when adequate dietary supplementation is not practical.

Hydralazine, with or without thiazides, may have shown some nitrogen retention. It seems likely that this was caused indirectly by the lowering of the blood pressure, which in turn reduced renal blood flow. If a patient with a history of renal insufficiency or renal insufficiency is observed, it may be desirable to discontinue use of hydrochlorothiazide.

In patients with renal disease, thiazides may be especially dangerous. Clinical evidence of renal failure may develop in patients with impaired renal function.

Dosage should always be carefully titrated. Pay special attention to the electrolyte balance of patients receiving hydralazine. In patients with cirrhosis and ascites, thiazides have produced symptoms of impending hepatic coma: confusion, drowsiness, tremor, laboratory tests revealed increased serum bilirubin, increased serum transaminases, and increased sodium and potassium excretion.

Thiazides derivatives, particularly in large doses, may decrease arterial responsiveness; therefore, hydralazine should be used with caution.

Hyperuricemia, occasionally with gout, may occur in patients receiving hydrochlorothiazide. The hyperuricemia is generally reversible and the serum uric acid level is usually normal.

Thiazides may decrease arterial responsiveness to tubocurarine; if possible, withdraw therapy two weeks prior to surgery. If emergency surgery is required, thiazides have been observed. If emergency surgery is indicated, preanesthetic and anesthetic agents should be administered in reduced dosage.

Thiazide sensitivity reactions should be considered in patients with a history of allergy or bronchial asthma.

**Usage in Pregnancy**

The safety of reserpine for use during pregnancy or lactation has not been established; therefore, the drug should be used in pregnant patients or in women of childbearing age only when the benefit to the patient, in the judgment of the physician, is essential to the welfare of the patient. Increased respiratory tract secretions, nasal congestion, coryza, and anorexia may occur in pregnant patients. The drug may cross the placental barrier and is known to cross the placental barrier, and to appear in breast milk.

Hyperuricemia

Although there has been no adverse experience with hydralazine in pregnancy, this drug should be used in pregnancy only when, in the judgment of the physician, it is deemed essential to the welfare of the patient.

**Hydralazine**

Thiazides should be used with caution in pregnant or lactating patients, since they cross the placental barrier and may result in fetal hyperbilirubinemia, thrombocytopenia, or altered carbohydrate metabolism. It is therefore possible that adverse reactions seen in the adult may occur in the newborn.

### PRECAUTIONS

Reserpine increases gastrointestinal motility and secretion. It should be used cautiously in patients with a history of peptic ulcer, ulcerative colitis, or other gastrointestinal disorders. It may precipitate hypoglycemia in diabetics.

Because of the effect of catecholamine depletion, asthmatics are more apt to be hypersensitive to the drug and their condition may be aggravated. Therefore, their condition may be aggravated.

Hydralazine

The following adverse reactions have been associated with the use of thiazide diuretics:

adjust poorly to lowered blood pressure levels. Use reserpine cautiously with digitalis and quinidine since cardiac arrhythmias have occurred with reserpine. Concurrent use of guanethidine and rauwolfia derivatives may cause bradycardia, mental depression, and postural hypotension.

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Preoperative withdrawal of reserpine does not assure that circulatory instability will occur. It is important to withdraw reserpine at least two weeks before surgery and consider this in the overall management, since hypertension has occurred in patients receiving rauwolfia derivatives. Antihypertensive agents (e.g., reserpine, guanethidine, etc., etc., norepinephrine) have been employed to treat severe hypovascular effects.

Hydralazine stimulation produced by hydralazine can cause anginal attacks and ECG changes of myocardial ischemia. The drug has been implicated in the production of myocardial infarction. It must, therefore, be used with caution in patients with suspected coronary artery disease.

The "hyperdynamic" circulation caused by hydralazine may accentuate special cardiovascular hazards, particularly in patients with a history of increased pulmonary artery pressure in patients with mitral valvular disease. The drug may reduce the pressor responses to epinephrine. Postural hypotension may result from hydralazine but is less common than with reserpine. Drug-induced depression may persist for several months after drug withdrawal.

Hydralazine should be discontinued for at least two weeks before giving electroshock therapy. MAO inhibitors should be avoided or used with extreme caution.

Hydralazine administration of doses over 400 mg per day may produce in a few patients an arthritic-like syndrome leading to a clinical picture simulating acute systemic lupus erythematosus. A rare but serious reaction may occur at lower doses and symptoms and signs usually regress when the drug is discontinued, but long-term treatment with steroids may be necessary.

Hydralazine may be associated with a history of hepatic coma. L.E. cells may be found in the blood of patients on the drug who are asymptomatic. An L.E. cell preparation is indicated if the patient has arthralgia, fever, testicular pain, continued malaise, or other unexplained symptoms.

Use MAO inhibitors with caution in patients receiving hydralazine. **Hydralazine**

There have been several reports, published and unpublished, concerning nonspecific small bowel lesions consisting of stenosis, with or without ulceration, associated with the administration of enteric-coated hydralazine. These lesions may occur with plain tablets or with coated tablets. These lesions may occur with enteric-coated potassium tablets alone or when they are used with nonenteric-coated thiazides for certain other oral diuretics.

These lesions have caused obstruction, hemorrhage, and perforation. Surgery was frequently required and deaths have occurred.

Available information tends to implicate enteric-coated hydralazine. These lesions may occur with plain tablets or with coated tablets. These lesions may occur with enteric-coated potassium tablets alone or when they are used with nonenteric-coated thiazides for certain other oral diuretics.

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### Guanethidine

As with all antihypertensive agents, give cautiously to patients with severe coronary insufficiency, recent myocardial infarction, and other cardiovascular reactions.

**Hematologic:** Leukopenia, thrombocytopenia, agranulocytosis, aplastic anemia.

**Cardiovascular:** Orthostatic hypotension may occur and may be potentiated by alcohol, barbiturates, or narcotics.

**Miscellaneous:** Muscle spasm, weakness, restlessness. Whenever adverse reactions are moderate or severe, the dose should be reduced or therapy withdrawn.

**DOSE AND ADMINISTRATION**

One or 2 tablets daily. To initiate therapy, 1 tablet daily for 1 to 2 weeks.

Since the antihypertensive effects of reserpine are not immediately apparent, maximal reduction in blood pressure from a given dose of Ser-Ap-Es may not be attained until the second week. The dosage should be increased by at least 50 per cent. Watch effects carefully.

**HOW SUPPLIED**

Tablets (dark salmon pink, dry-coated), each containing 10 mg reserpine, 25 mg hydralazine hydrochloride, and 15 mg hydrochlorothiazide; bottles of 100 and 1000.

Rev. 3/72

## Esimil®

guanethidine monosulfate 10 mg  
hydrochlorothiazide 25 mg

### INDICATIONS

Esimil is indicated for all cases of hypertension, except the mildest and the most severe.

### CONTRAINDICATIONS

Known hypersensitivity, mental depression (especially with suicidal tendencies), active peptic ulcer, ulcerative colitis, digitalis intoxication, edema, pregnancy, and patients receiving electroconvulsive therapy, hydralazine.

Hypertension in patients in general have a higher risk of intrapartum hypertension and other cardiovascular complications than normotensive patients.

Reserpine-treated patients are not known to have a higher risk of hypertension than those with otherwise comparable hypertensive patients.

Preoperative withdrawal of reserpine does not assure that circulatory instability will occur.

It is important to withdraw reserpine at least two weeks before surgery and consider this in the overall management, since hypertension has occurred in patients receiving rauwolfia derivatives.

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Preoperative withdrawal of reserpine

## driving me nuts...

continued from page 17

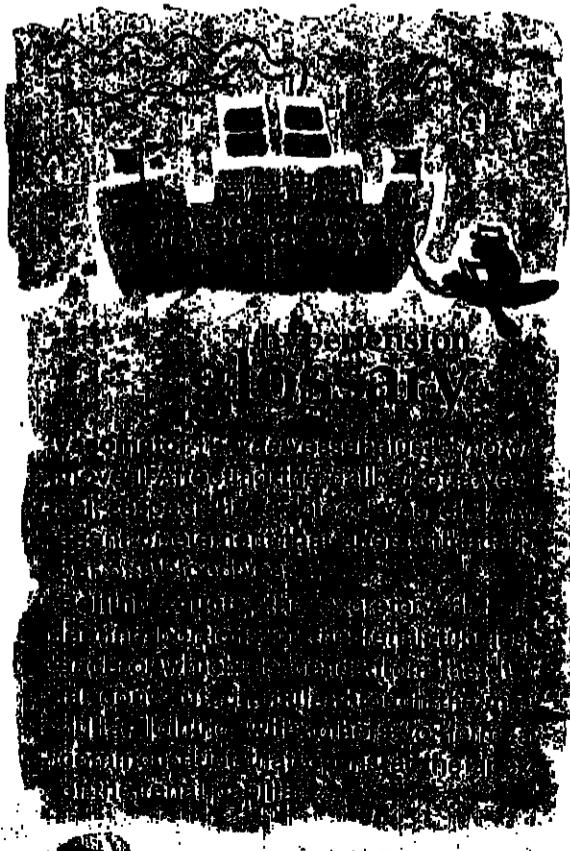
light. Their response determines whether or not they will get an electric shock. Needless to say, the situation is totally foreign to these very gregarious, normally active primates, who generally run in groups of about 20 in the wild.

"It is quite possible to make a human analogy here. Most of us have to curtail our gregariousness each day at work, confining ourselves to the small space of an office, or a laboratory, or a position on an assembly line. We, too, respond to cues, although they are much more subtle and complex. We respond to the alarm clock, the telephone, the lunch whistle. If we do not, there are noxious stimuli—the displeasure of superiors, no promotion, the annoyance of fellow workers."

### Catheters worn continuously

In two groups of monkeys, implanted aortic catheters continuously record blood pressure levels associated with environmental cues. Preliminarily, both groups of animals are subjected to exactly the same treatment, and they show approximately the same blood pressures. Both groups wear their catheters continuously, 24 hours a day. Both live in isolation booths for a period of each day.

The change comes after this preliminary training period. In the experimental group of 11 animals, each is conditioned to press a switch key whenever a light goes on, because he learns that if he fails to switch the light off he will get an electric shock. His heart rate and arterial blood pressure increase as he goes for the key, and in a few weeks the mean pressure rise reaches 20 mm. Hg. When the animal switches the light off its blood pressure returns to base levels. But after a few months of being powerfully and continuously conditioned by environmental stimuli, an animal's mean arterial blood



pressure elevation begins to persist between daily sessions. Seven control monkeys—not subjected to flashing lights and shock—have had no rises in pressure.

High pressure levels in the experimental monkeys have peaks and valleys. On days set aside for behavioral studies, arterial blood pressure is highest in the isolation booth. Afterwards, blood pressure declines to slightly lower levels. The biggest drop in arterial pressures is recorded immediately after the animal is removed from the isolation chamber. After this period of relief, the pressure gradually rises, up to the time of the next daily session in the lights cage, when it takes another spurt. From these slowly rising pressure values, it appears that the animal foresees each day's session with the lights.

To assess whether the muscular act of pressing the key raised the animal's blood pressure, the key is removed from the apparatus. The light flashes as before, and only an animal's deliberate, self-determined rise in pressure foretells delivery of shock stimuli. The animals have soon learned to raise their blood pressure in response to the lights.

Here, too, Dr. Herd makes some cautious human analogies. "Perhaps this happens in our culture. Maybe we are rewarded not so much for performing the task, but for being crisp and responsive—or 'revved up'—in anticipation of the task. Our society tends to reward people who are aggressive, outgoing, brisk. Certainty and authority are very highly regarded."

### Analogy to humans

Comparisons between human beings and squirrel monkeys are safely made on physiologic grounds, according to Dr. Herd. "Both species of primates have identical organs. So far as we know, their organs work in the same way, with similar hormone responses of adrenal cortical steroids and adrenal medullary secretions.

"But there are some differences. Size is the most obvious. The squirrel monkey is about a foot long, and weighs less than 2 pounds. Size differences account for metabolic differences. All small primates have a higher metabolic rate than man and a slightly higher resting blood pressure."

This higher metabolic rate, says Dr. Herd, makes the squirrel monkey more typical of a particular group of people than of all people. "These monkeys are more like labile hypertensives encountered in clinical medicine than any other creature we have found. They're susceptible to a high-fat diet, and they develop hardening of the arteries just as humans do. Atherosclerotic changes in their blood vessels are microscopically indistinguishable from those in humans. So are biochemical and pathological changes. Other animals—including dogs, rabbits, rats, and guinea pigs—get hardening of the arteries, but they show different lesions in their blood vessels."

"In the lab, we feed our healthy monkeys a diet with the same composition of proteins, carbohydrates, and fats recommended for healthy humans."

But the most striking similarity between the hypertension of human beings and squirrel monkeys comes from Dr. Herd's experimental data: not all mon-

keys develop hypertension under pressure. Only nine out of 11 experimental animals did. Therefore, whether monkey or human, some individuals are more susceptible of hypertension in their environment than others.

This fact is reflected in statistics showing that some hard-driving executives who thoroughly enjoy their jobs are just as likely to get hypertension as their driven employees. Dr. Peter B. Dews, who was trained as a physician and surgeon at the University of Leeds in England, and who is now Stanley Cobb Professor of Psychiatry and Psychobiology in Harvard's Department of Psychiatry, says:

"It may turn out that it does not matter how blood pressure is raised—whether by pleasure or non-pleasure. It may be that the mere act of raising the pressure is



what produces human hypertension. Perhaps repeated cumulative periods of high blood pressure over a period of time will do it. If so, there may be some value in searching for new prophylactic, blood pressure-lowering drugs that could be given before stressful situations develop."

One way of lowering blood pressure—in the laboratory, at least—has already been found by the Harvard group. They teach squirrel monkeys to lower blood pressure in much the same way they taught them to raise it.

Does this have any human application?

"I really don't know," says Dr. Dews.

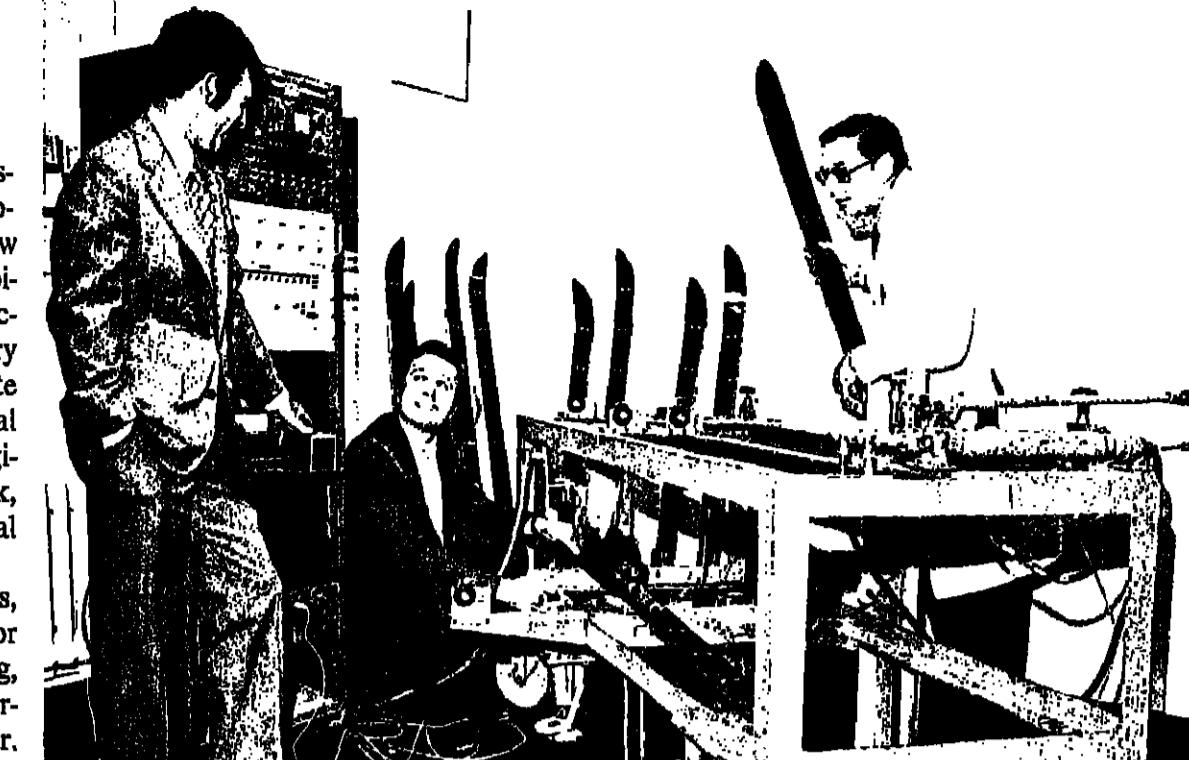
Human studies are scheduled to begin shortly at Massachusetts General Hospital under the direction of Dr. Edgard Haber, Professor of Medicine at Harvard. These studies will be grounded in the primate data accumulated thus far, plus a hint found recently in Cannon's handwritten diary: that pathologic effects of emotion may be due to failure to have normal exit in muscular movement.

## Sports-Related Injuries Are Focus of Youth Unit

TREATMENT of sports-related injuries in adolescents is the prime concern of the recently established Rainbow Sports Medicine Center at Rainbow Children's Hospital, part of the University Hospitals of Cleveland. Training, research into the effectiveness of sports equipment, and methods of injury treatment and prevention in the high school athlete are other activities studied at the center, an unusual combination of medical school, hospital, and engineering school, according to Dr. Robert Mack, head of orthopedic surgery at Cleveland General Hospital and director of the center.

The center employs the science of biomechanics, the application of mechanical laws to the locomotor system, in studying the body's reactions to padding, methods of taping, equipment, and playing surfaces. Heading the biomechanical studies is Dr. Victor Frankel, director of research at the facility.

One of the courses offered by the center is for nonplaying students who participate in school athletic programs as managers and junior trainers. They learn techniques of training, exercise, and taping, allowing them a greater role in assisting their coaches and trainers. Working with the center in an advisory capacity is a board made up of Cleveland-area educators associated with athletics from the high school to the college level.



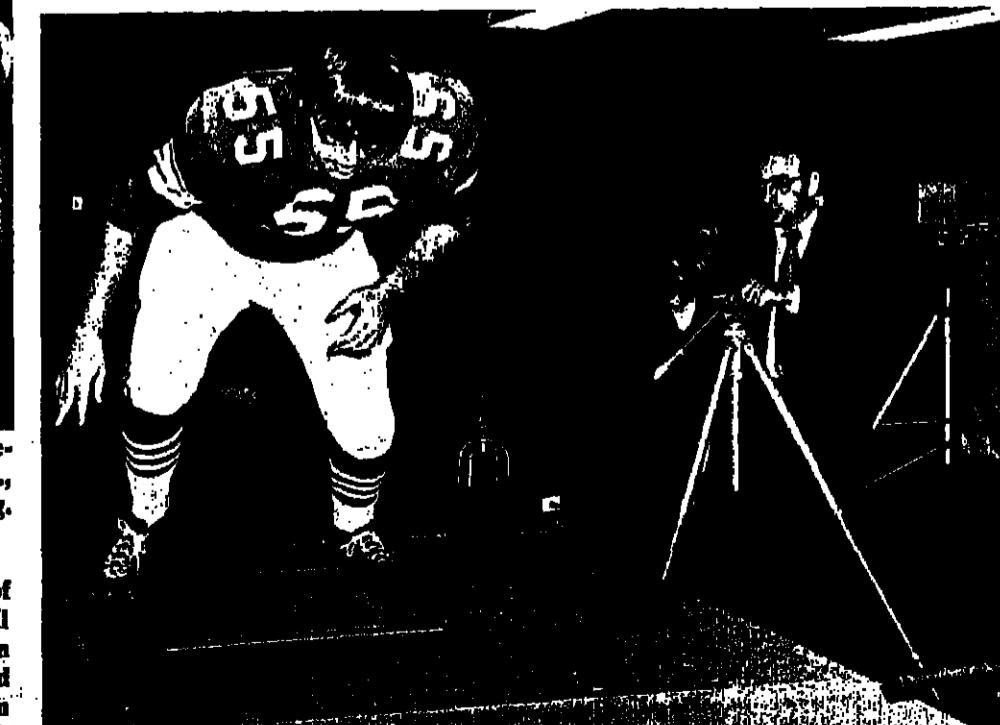
Staff members (left to right) Dr. Mack, Eugene Bahnuk, Ph.D., and Dr. Frankel demonstrating the apparatus that tests ski bindings. The center's study of the failure of most bindings to protect the skier won awards from the U.S. Ski Association and from the American Academy of Orthopedic Surgeons.



Young athlete, above, has his coordination tested by Dr. Frankel, who designed the testing device at the biomechanics lab at Case Western Reserve U., where he is Professor of Orthopedic Surgery and Biomedical Engineering.



Goalie for the Cleveland Crusaders hockey team has his arm checked by Dr. Mack, team physician. Some staff members are connected with Olympics.



Albert Burstein, D.S.M.E., of the center, photographs football player in motion. Athlete is on force-plate, a device that is used to measure the ground reaction force of the runner's take-off.

# Keeping the mild hypertensive in his place

## that's "Antihypertenacity" Esidrix has it (hydrochlorothiazide)

Esidrix not only gets blood pressure down, and gets it down smoothly, but it keeps on exerting its antihypertensive effect.

Still unsurpassed as a basic diuretic-antihypertensive, Esidrix has the gradual, sustained action needed in the long-term management of mild hypertension. We call it antihypertenacity.

And as a diuretic, Esidrix is useful in many forms of edema.

Contraindications include anuria. Use with caution in patients with impaired renal or hepatic function.



### Esidrix® (hydrochlorothiazide)

Indications: Hypertension and edema.

Contraindications: Anuria; hypersensitivity to this or other sulfonamide-derived drugs. The routine use of diuretics in an otherwise healthy pregnant woman with or without mild edema is contraindicated and possibly hazardous.

Warnings: Use with caution in severe renal disease. In patients with renal disease, thiazides may precipitate azotemia. Cumulative effects of the drug may develop in patients with impaired renal function.

Thiazides should be used with caution in patients with impaired hepatic function or progressive liver disease, since minor alterations of fluid and electrolyte imbalance may precipitate hepatic coma.

Thiazides may be additive or potentiative of the action of other antihypertensive drugs. Potentiation occurs with ganglionic or peripheral adrenergic blocking drugs.

Sensitivity reactions may occur in patients with a history of allergy or bronchial asthma.

The possibility of exacerbation or activation of systemic lupus erythematosus has been reported.

### Usage in Pregnancy

Usage of thiazides in women of childbearing age requires that the potential benefits of the drug be weighed against its possible hazards to the fetus. The hazards include fetal or neonatal jaundice, thrombocytopenia, and possibly other adverse reactions which have occurred in the adult.

### Nursing Mothers

Thiazides cross the placental barrier and appear in cord blood and breast milk.

Precautions: Periodic determination of serum electrolytes to detect possible electrolyte imbalance should be performed at appropriate intervals. Observe patients for clinical signs of fluid or electrolyte imbalance (hyponatremia, hypochloremic alkalosis, and hypokalemia). Serum and urine electrolyte determinations are particularly important when the patient is vomiting

excessively or receiving parenteral fluids. Medication such as digitalis may also influence serum electrolytes. Warning signs are dryness of mouth, thirst, weakness, lethargy, drowsiness, restlessness, muscle pains or cramps, muscular fatigue, hypotension, oliguria, tachycardia, and gastrointestinal disturbance such as nausea or vomiting.

Thiazides may develop with thiazides as with any other potent diuretic, especially during brisk diuresis, when severe cholestasis is present, or during concomitant administration of steroids or ACTH.

Interference with adequate oral intake of electrolytes may also contribute to hypokalemia. Diuretics may exaggerate metabolic effects of other hypotensive agents especially with reference to myocardial activity.

Any chloride deficit is generally mild and usually does not require specific treatment except under extraordinary circumstances (as in liver disease or renal disease).

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BEHIND EACH PRODUCT  
A TRADITION OF  
BASIC RESEARCH

### How much drug to prescribe?

How well is it working?

CIBA-GEIGY scientists help find the answers. Their new analytical methods detect blood levels as low as 0.05 micrograms per milliliter, thus helping establish proper dosage ranges.

Our contribution to medicine goes far beyond producing it.

CIBA

## One Man... and Medicine

ARTHUR M. SACKLER, M.D.,  
International Publisher, Medical Tribune



### Is That Test Necessary?

THE FOUR DOCTORS at dinner one night had just finished when the youngest said, "Boy, did I get chewed out today! I didn't have a spinal tap on one of my night admissions by morning rounds."

"Are you still doing your own lab work-up at night?"

"Of course, and what a waste. It really doesn't make any sense at all. Why does every patient have to have, in addition to physical, history, urine and blood work-up, virtually automatic ECGs, x-rays, and—on the basis of a remote differential diagnosis—spinal taps?"

### Status Medicine

"I've done a stint in one of the African countries which is really short in medical manpower. We were lucky to be able to do microscopies on urine, blood, and stool. There were more important things to do all the time. This is status medicine," the young doctor said. "It makes the doctor feel good and the hospital look good, but, in the over-all view, how much does it really contribute to the patient? Shouldn't my chief's question have been, Why did you do that spinal tap or the ECG or that x-ray, and not the other way around? What's 'good' in a teaching hospital could be considered 'economic exploitation' of the patient in private practice."

If you ever had a post-spinal tap headache and tinnitus and had it drag on for months, you wouldn't throw spinal taps around. I couldn't help remembering that over three decades ago it was considered good medicine in the hospital where I interned to do x-ray polimetry on every pregnant woman. I shudder to think of the fetal and genetic damage that these routines of 'scientific' or advanced medical care produced. How many things are we doing today that are comparable?

Some time ago, in Europe, I was shown a miraculous urine analysis machine by its breathless promoter. This first "autoanalyzer" I was told, could do a thousand urine analyses a day, but the next generation machine they were going to build would be able to do 10,000 urines a day. In growing astonishment I exclaimed, "Where on earth are you going to get 10,000 urine specimens a day?" And then, too, a friend to whom I told this story remarked, "And what are they going to do the day after?"

Which, of course, brings us around to the fundamental thing we have observed before—what this country needs and maybe what the world needs is not just "a good 5¢ cigar," but a lot more good, old-fashioned clinical sense and clinical medicine.

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### Cancer Unit Prepares Child for Home Life



## 2 More Doctor Units Sign Up As Unionizing Trend Grows

Continued from page 1

the county of its medical services with social service and welfare agencies.

"One of our complaints," Dr. J. Lee Aiken, president of the physicians' group, told MEDICAL TRIBUNE, "was that the county board of supervisors tried to put the county hospital under the welfare director. Furthermore, they refused to talk to the medical staff regarding patient care and hospital administration. We felt that the physicians should have some input into decisions affecting medical services."

### Economics and Economy

Curiosity got the better of me. How much will it cost for the individual patient going through the screen?

"Oh," the answer was, "\$60 to \$70."

But can you sell enough of these integrated units to get into mass production, assuming that governmental agencies would want so complex a screen procedure?

"Sure, we think we can sell it to Latin-American governments interested in health."

Can you? Do you realize what percentage of the populations in some of the countries have sections of their economy with a per capita gross national product of \$100 to \$200 annually?

### Now, Improved Machine and Reality

There was a blank look. Could it be that so simple a fact can be obscured by the beauty and glamour of our technologic intricacies?

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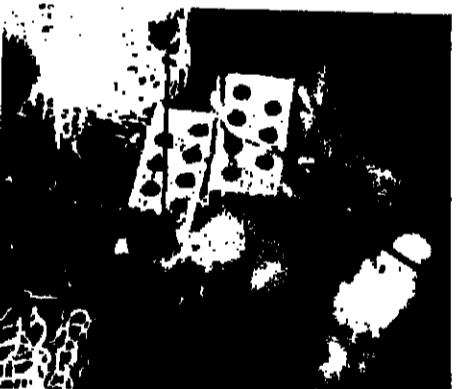
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# Extending the boundaries of knowledge in modern brain research



## Remote-control ESB:

In experiments by Delgado and associates, electrodes are implanted into specific brain areas preparatory to behavior programming by remote-control electrostimulation of the brain.



## Radio-controlled ESB pinpoints action of Librium (chlordiazepoxide HCl) on selected brain areas of rhesus monkeys

Remote-control ESB (electrostimulation of the brain) elicited predictable behavior patterns in monkeys, patterns that persisted only as long as the specific stimulation was applied. Librium was then administered to determine its effect on the ESB-altered behavior patterns. Delgado and associates,<sup>1,2</sup> working with Librium, have helped to elucidate the CNS action of this psychotropic agent in monkeys.

Experimental observations<sup>1,2</sup> in monkeys\* showed that:

- Librium (chlordiazepoxide HCl) blocked an electrically stimulated epileptogenic response of the amygdala, including the occurrence of an "after-discharge." Hostility of the monkey was controlled.

- Librium reduced the excitability of the monkey's central gray area, a brain structure apparently related to aggressive behavior and pain perception.
- Librium did not modify the appetite-inhibiting effects of caudate nucleus stimulation.
- Librium did not change the motor effect of internal capsule stimulation, which produced flexion of the monkey's arm and leg.
- Librium also decreased total activity in gibbons but favored normal activity such as grooming and play.

1. Delgado, J. M. R.; Bracchitta, H., and Snyder, D. R.: "Psychoactive Drugs and Radio-Controlled Behavior," film presented at the 124th Annual Meeting, American Psychiatric Association, Washington, D.C., May 3-6, 1971.
2. Delgado, J. M. R., et al.: "Radio Communication with the Brain," Scientific Exhibit presented at the 124th Annual Meeting, American Psychiatric Association, Washington, D.C., May 3-6, 1971.

\*While the animal experiments described can be used to obtain a better understanding of the action of Librium (chlordiazepoxide HCl) in monkeys, no clinical conclusions can be drawn, as it is not possible to extrapolate animal data to humans.

Specific calming action in monkeys indicated in experimental studies

**Librium®**  
(chlordiazepoxide HCl)

## Clinical experience with Librium® (chlordiazepoxide HCl)

After more than 12 years of wide clinical use, experience with Librium (chlordiazepoxide HCl) continues to reflect its favorable therapeutic index. By its antianxiety action, Librium can help encourage activity of ambulatory patients with deleterious anxiety and can enhance their participation in productive, recreational or rehabilitative activities.

On proper maintenance dosage, Librium generally helps calm the patient, usually without unduly interfering with mental acuity or ability to perform. When excessive anxiety has been reduced to appropriate levels, Librium therapy should be terminated.

Librium is used concomitantly with certain specific medications of other classes of drugs, such as cardiac glycosides, diuretics and antihypertensive agents, whenever anxiety is a clinically significant factor.

Before prescribing, please consult complete product information, a summary of which follows:

**Indications:** Relief of anxiety and tension occurring alone or accompanying various disease states.

**Contraindications:** Patients with known hypersensitivity to the drug.

**Warnings:** Caution patients about possible combined effects with alcohol and other CNS depressants. As with all CNS-acting drugs, caution patients against hazardous occupations requiring complete mental alertness (e.g., operating machinery, driving). Though physical and psychological dependence have rarely been reported on recommended doses, use caution in administering to addiction-prone individuals or those who might increase dosage; withdrawal symptoms (including convulsions), following discontinuation of the drug and similar to those seen with barbiturates, have been reported. Use of any drug in pregnancy, lactation, or in women of childbearing age requires that its potential benefits be weighed against its possible hazards.

**Precautions:** In the elderly and debilitated, and in children over six, limit to smallest effective dosage (initially 10 mg or less per day) to preclude ataxia or oversedation, increasing gradually as needed and tolerated. Not recommended in children under six. Though generally not recommended, if combination therapy with other psychotropics seems indicated, carefully consider individual pharmacologic effects, particularly in use of potentiating drugs such as MAO inhibitors and phenothiazines. Observe usual precautions in presence of impaired renal or hepatic function. Paradoxical reactions (e.g., excitement, stimulation and acute rage) have been reported in psychiatric patients and hyperactive aggressive children. Employ usual precautions in treatment of anxiety states with evidence of impending depression; suicidal tendencies may be present and protective measures necessary. Variable effects on blood coagulation have been reported very rarely in patients receiving the drug and oral anticoagulants; causal relationship has not been established clinically.

**Adverse Reactions:** Drowsiness, ataxia and confusion may occur, especially in the elderly and debilitated. These are reversible in most instances by proper dosage adjustment, but are also occasionally observed at the lower dosage ranges. In a few instances syncope has been reported. Also encountered are isolated instances of skin eruptions, edema, minor menstrual irregularities, nausea and constipation, extrapyramidal symptoms, increased and decreased libido—all infrequent and generally controlled with dosage reduction; changes in EEG patterns (low-voltage fast activity) may appear during and after treatment; blood dyscrasias (including agranulocytosis), jaundice and hepatic dysfunction have

been reported occasionally, making periodic blood counts and liver function tests advisable during protracted therapy.

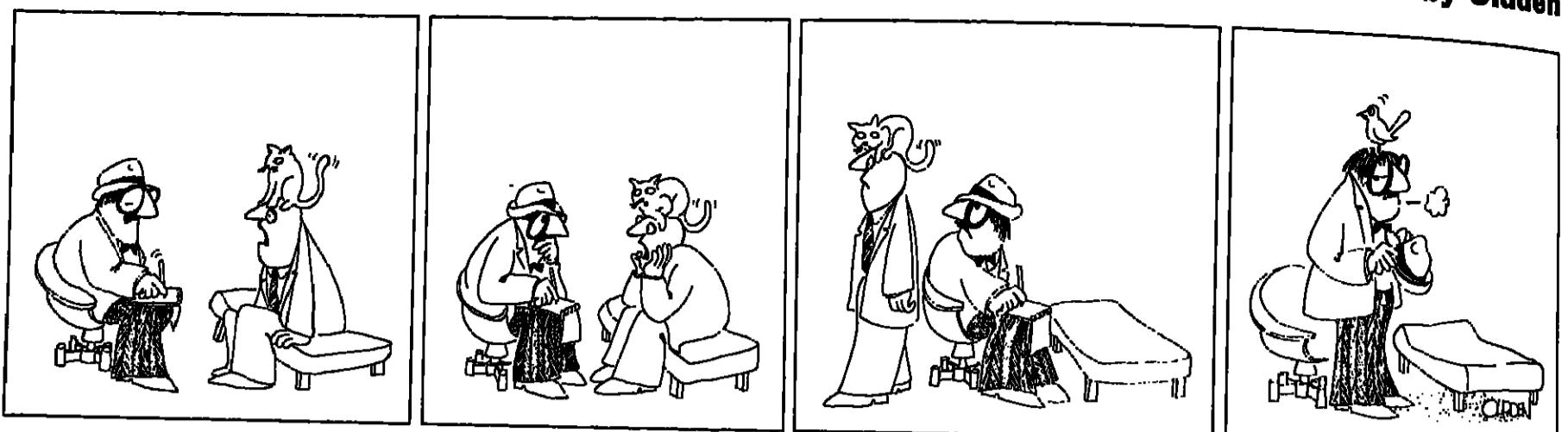
**Supplied:** Librium® Capsules containing 5 mg, 10 mg or 25 mg chlordiazepoxide HCl. Libritabs® Tablets containing 5 mg, 10 mg or 25 mg chlordiazepoxide.

for the relief of clinically significant anxiety in emotional and somatic disorders: a wide range of dosage options

**Librium®**  
(chlordiazepoxide HCl)  
5-mg, 10-mg, 25-mg capsules  
up to 100 mg daily  
in severe anxiety



Roche Laboratories  
Division of Hoffmann-La Roche Inc.  
Nutley, N.J. 07110



### SURGICAL NOTES

#### Surgery for Tennis Elbow

LAS VEGAS, Nev.—Tennis elbow is most often cured by rest or hormonal injections, but occasionally surgery is necessary.

Dr. Harold B. Boyd, Emeritus Professor of Orthopaedic Surgery at the University of Tennessee, said that of 871 tennis-elbow patients seen at the Campbell Clinic in Memphis over a 16-year period, only 40 did not respond to the conservative treatment and required surgery. In four patients, bilateral operations were performed.

The surgery brings relief of pain and restoration of full range of motion in almost all cases, said Dr. Boyd. The patient requires three to six months to regain full strength in the forearm. Average time for returning to work and hobbies was six weeks.

Speaking to the annual meeting of the American Academy of Orthopaedic Surgeons here, Dr. Boyd remarked that the arm is placed in a sling postoperatively, but active motion is started in 24 hours.

He remarked that probably most tennis-elbow patients are never seen by a doctor. Healing by conservative treatment usually occurs within six months, he said, and recurrences of the disorder are rare—only about 3 per cent.

Conauthor was Dr. Andin C. McLeod, Jr., of Hattiesburg, Miss.

#### Thromboembolic Snags

STOCKHOLM—Thromboembolic complications in major surgical interventions still constitute a serious problem, but recent studies have shown that there are possibilities of reducing their frequency, according to an editorial in a recent issue of the *Journal of the Swedish Medical Association*.

One study, it said, demonstrated that a small dose of heparin subcutaneously before and for a week after operation reduces the incidence of venothrombosis from 42 per cent to 8 per cent. Another indicated that three doses of heparin prevent post-operative thrombosis after major abdominal intervention for benign disorders just as effectively as prolonged subcutaneous heparin prophylaxis. Still another study found that dextran administered in con-

nection with surgery reduces the thrombosis frequency by one-half in many patients.

Such preventive methods appear to be superior to therapeutic exercise or early ambulation, but before a definite stand is taken on routine prophylaxis with either heparin or dextran, it would be desirable to see the results of long-term studies on representative material, the editorial said.

#### High Blood Pressure

STOCKHOLM—Results with baropacing, the stimulation of the sinus nerve, in five patients with therapy-resistant severe essential hypertension were reported by Dr. Lennart Hansson, of the University of Michigan Medical Center, at the annual meeting of the Swedish Medical Society.

Electrodes were implanted bilaterally around the sinus nerve and connected to a Medtronic baropacer placed subcut-

taneously in the region of the pectoralis. Stimulation was aided by an external radiofrequency transmitter.

Dr. Hansson and his associates, Drs. Calvin Ernst, Stephen H. Hunyor, and Steve Julius, observed, at the onset of stimulation, a rapid drop in median arterial pressure of 22 mm. Hg. The cardiac index and heart frequency were influenced only insignificantly. Peripheral vessel resistance sank by 19 per cent.

normal; this was associated with an end-diastolic pressure of 21 mm. Hg, in contrast to 9 mm. Hg in the normals. There was also a marked reduction in the mean rate of fiber shortening during ejection in the hypertensives, "which resulted in a profound reduction in ejection fraction, and because of this subnormal emptying, end-systolic volume was approximately twice normal."

#### Significant Impairment Demonstrated

Measures of the contractility of the myocardium demonstrated a significant impairment in the hypertensive group, the investigators said.

When the hypertensives were subjected, by leg elevation, to a 10 per cent rise in ventricular end-diastolic volume, they reported, "the normal increase in ejection fraction and stroke volume did not occur."

As a result, end-systolic volume rose significantly, "indicating inadequate emptying in response to the stress of acutely increased preload." Moreover, when they were subjected, by sustained hand grip, to a significant increase in aortic pressures, "ejection fraction and stroke volume fell."

It was noted that the patients were operating with a preload 30 per cent larger than

volume, "demonstrating inadequate emptying in response to the stress of acutely increased afterload."

All these results were said to reflect impaired contractility.

Their study, the investigators declared, "demonstrates that even without the classical symptoms and signs of decompensation, contractile element failure in hypertensive heart disease can be identified by a simple noninvasive test—that is, the chest x-ray."

The authors were Drs. Ernesto Rodriguez, Ravinder Narang, E. Sultan Ahmed, James J. Fiore, and Gilbert E. Levinson.

#### Spina Bifida Group Forms

Medical Tribune Report

CHICAGO—The Spina Bifida Association of America was formed here recently at a meeting of 80 delegates from 27 organizations representing more than 3,000 patients with spina bifida. It will seek, among other objectives, to create a better understanding of the problems of persons with this defect. An estimated 11,000 infants are born with spina bifida each year.

#### Lead Poisoning

NEWARK, N.J.—Progress in the fight to wipe out lead poisoning among children in this community has been made evident through a study of hospital admission records, according to Dr. Ann Browder, Dr. Donald B. Louria, and Morris Joselow, Ph.D., of the New Jersey Medical School.

They said that the admissions data reflected the efforts of an intensified blood-screening program started in 1969 with the development of an environmental toxicology unit of the college, working in collaboration with the Newark Department of Health and Welfare and the State Department of Health.

The analysis of hospital records showed a marked reduction in average blood-lead levels—from 130 to 86 micrograms per 100 ml.—in asymptomatic children. Intensified screening also produced about six times as many hospital admissions in 1970 (18.2 a month) as in 1967-68 (3.2 a month), mainly because many more children were being tested and treated for lead poisoning, the study found.

#### Sudden Death Syndrome

ADELAIDE, AUSTRALIA—Sudden death syndrome, or "cot death," has become a major contributory cause of infant mortality in South Australia, and in the age group two to seven months it now accounts for 60 per cent of all deaths, a survey here showed.

In children aged two weeks to two years, it leads the list of mortality causes, ahead of congenital malformation, infections, and accidents, said Dr. Susan Beal, a pathologist at Adelaide Children's Hospital.

#### Diagnosis of Hemophilia

ULM, WEST GERMANY—Early diagnosis can help increase the life expectancy of hemophiliacs, participants at the annual congress of the Hemophilic Association of Germany were told.

Dr. M. H. Maurer, president of the association, noted that life expectancy has increased from 15 to 40 and even 50 years with modern treatment.

The congress called for a network of treatment centers throughout West Germany to help the nation's 30,000 hemophiliacs.

#### Nutritional Anemia in India

NEW DELHI—One child in two in India's population suffers from nutritional anemia, according to a survey by the Indian Council of Medical Research in association with state nutrition centers.

The survey also showed that about 50,000,000 children one to six years old are affected by protein-calorie malnutrition.

From animal studies to clinical studies to sleep research laboratory studies in man...

### Multiphasic testing documents the effectiveness and relative safety of Dalmame (flurazepam HCl) for sleep

One 30-mg capsule h.s.—usual adult dosage. One 15-mg capsule h.s.—initial dosage for elderly or debilitated patients.

Before prescribing Dalmame (flurazepam HCl), please consult Complete Product Information, a summary of which follows.

**Indications:** Effective in all types of insomnia characterized by difficulty in falling asleep, frequent nocturnal awakenings and/or early morning awakening. In patients with recurring insomnia, poor sleeping habits; and in acute or chronic medical situations requiring restful sleep. Since insomnia often transient and intermittent, prolonged administration is generally not necessary or recommended.

#### Contraindications: Known hypersensitivity to flurazepam HCl.

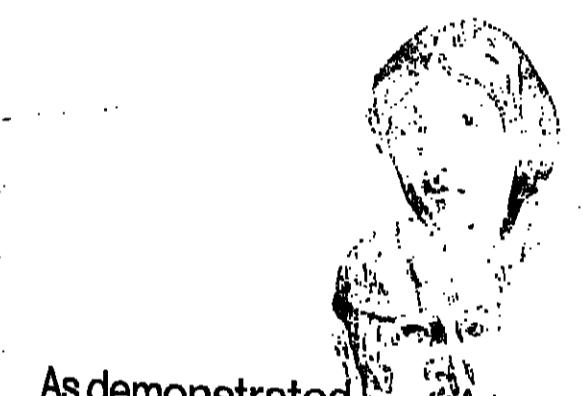
**Warnings:** Caution patients about possible impaired effects with alcohol and other CNS depressants. Caution against hazardous occupations requiring complete mental alertness (e.g., operating machinery, driving). Use in women who are or may become pregnant only when potential benefits have been weighed against possible hazards. Not recommended for use in persons under 15 years of age. Though physical and psychological dependence

have not been reported on recommended doses, use caution in administering to addiction-prone individuals or those who might increase dosage. **Precautions:** In elderly and debilitated, initial dosage should be limited to 15 mg to produce oversedation. Dizziness, drowsiness and/or confusion combined with other drugs having hypnotic or CNS depressant effects, consider potential additive effects. Employ usual precautions. Patients taking sedatives, lethargy, drowsiness and coma, probably indicative of drug tolerance or over dosage, have been reported. Also reported were headache, headache, hypertension, nausea, vomiting,

diarrhea, constipation, GI pain, nervousness, talkativeness, apprehension, irritability, weakness, palpitations, chest pains, body and joint pains and GU complaints. These have also been rare occurrences of sweating, flushing, difficulty in focusing, blurred vision, burning eyes, faintness, hypotension, shortness of breath, pruritus, skin rash, dry mouth, bitter taste, excessive salivation, anorexia, euphoria, depression, slurred speech, confusion, restlessness, hallucinations and elevated SGOT, SGPT, total and direct bilirubins and alkaline phosphatase. Paradoxical during repeated therapy. Observe special precautions in presence of impaired renal or hepatic function. **Adverse Reactions:** Dizziness, drowsiness, lightheadedness, staggering, confusion, hallucinations, flushing, particularly in elderly or debilitated patients. **Overdosage:** Sedation, lethargy, drowsiness and coma, probably indicative of drug tolerance or over dosage, have been reported. Also reported were headache, headache, hypertension, nausea, vomiting,

alkaline phosphatase. Paradoxical during repeated therapy. Observe special precautions in presence of impaired renal or hepatic function. **Overdosage:** Individual for maximum benefit effect. Adults: 30 mg usual dosage, 15 mg may suffice in some patients. **Elderly or debilitated patients:** 15 mg initially until response is determined. **Supplied:** Capsules containing 15 mg or 30 mg flurazepam HCl.

### As confirmed in sleep research laboratory studies

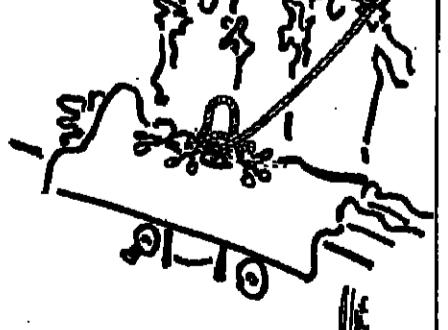
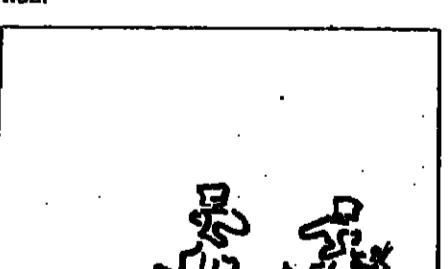


#### As demonstrated in clinical studies

• One 30-mg capsule of Dalmame (flurazepam HCl) at bedtime on average induced sleep within 17 minutes, decreased nocturnal awakenings, and provided 7 to 8 hours of sleep.

• Dalmame 30 mg was found to be effective for patients with difficulty in falling asleep, staying asleep or both.

• In studies to date, the effectiveness of Dalmame has been maintained without need to repeat or increase dosage.



Nerves like steel cables.

© 1970 Medical Tribune



## Diabetes Adjustment



Staff nurse, Linda Taylor gives patient individualized instruction, part of the diabetes adjustment program that is currently being offered at Creighton Memorial St. Joseph Hospital in Omaha.

now  
an ampicillin injection  
for routine office use.

## Polycillin® Intramuscular (sterile ampicillin trihydrate for suspension)

## Stability.

Polycillin Intramuscular is stable for 12 months as a dry powder. After reconstitution, it is stable for 60 days at room temperature.

## Convenience.

Stability facilitates routine use in office practice or on house calls...multi-dose vials allow reconstitution at your convenience, easily carried in your bag...ideal for initial therapy before a transfer to oral medication.

Economy. Stability permits use of multi-dose vials which substantially reduce the cost of delivering ampicillin by Intramuscular injection; each 10-cc. vial (2.5 Gm.) contains 10 doses of 250 mg. or 5 doses of 500 mg.

## BRIEF SUMMARY OF PRESCRIBING INFORMATION (13/1/72)

For complete information consult Official Package Circular.

Usage in Pregnancy: Safety for use in pregnancy is not established. Precautions: Mycotic or bacterial superinfection may occur. Cases of gonorrhoea with a suspected primary lesion of syphilis should have dark examinations before receiving treatment. In all other cases dark examinations should be suspended, monthly serological tests should be performed for a minimum of 4 months. Assess renal, hepatic, haemopoietic function and liver function during long-term therapy.

Adverse Reactions: Untoward reactions include: rash, urticaria, "hairy" tongue, nausea, vomiting and diarrhea, skin rash, urticaria, exfoliative dermatitis, erythema multiforme and anaphylaxis (usually with parenteral administration). Anemia, thrombocytopenia, thrombocytopenia purpura, eosinophilia, leukopenia, and agranulocytosis have been noted, are usually reversible and are believed to be hypersensitivity phenomena. Moderate elevations in SGOT have been noted.

Usual Dosage: Respiratory Tract Infections: Adults—250 mg. q.d.

Children—50 mg./Kg./day.

Gastrointestinal and Genitourinary Tract Infections: Adults—500 mg. q.i.d. Children—100 mg./Kg./day.

Urthritis in male adults due to *N. gonorrhoeae* 500 mg. b.i.d.

Children weighing more than 20 Kg. should be dosed according to the adult recommendations.

Contraindications: A history of allergic reactions to penicillin.

Warnings: Anaphylaxis may occur, particularly after parenteral administration and especially in patients with a history of allergies.

Penicillin or ampicillin reaction occurs, discontinue ampicillin.

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# the long-range analgesic

in chronic pain: continued relief without risk of tolerance

Though Talwin® Tablets can be compared to codeine in analgesic efficacy, Talwin is not subject to narcotic controls. For patients who require potent analgesia for prolonged periods, Talwin can provide consistent, long-range relief, with fewer of the consequences you've come to expect with narcotic analgesics.

- Comparable to codeine in analgesic efficacy: one 50 mg. Talwin Tablet appears equivalent in analgesic effect to 60 mg. (1 gr.) of codeine. Onset of significant analgesia usually occurs within 15 to 30 minutes. Analgesia is usually maintained for 3 hours or longer.
- Tolerance not a problem: tolerance to the analgesic effect of Talwin Tablets has not been reported, and no significant changes in clinical laboratory parameters attributable to the drug have been reported.
- Dependence rarely a problem: during three years of wide clinical use, only a few cases of dependence have been reported. In prescribing Talwin for chronic use, the physician should take precautions to avoid increases in dose by the patient and to prevent the use of the drug in anticipation of pain rather than for the relief of pain.
- Not subject to narcotic controls: convenient to prescribe—day or night—even by phone.
- Generally well tolerated by most patients: infrequently cause decrease in blood pressure or tachycardia; rarely cause respiratory depression or urinary retention; seldom cause diarrhea or constipation. If dizziness, lightheadedness, nausea or vomiting are encountered, these effects may decrease or disappear after the first few doses. (See next page of this advertisement for a complete discussion of Adverse Reactions and a Brief Summary of other Prescribing Information.)

50mg. Tablets

**Talwin®**  
brand of  
pentazocine  
(as hydrochloride)  
in moderate to severe pain

## in chronic pain: continued relief without risk of tolerance

Talwin® Tablets brand of pentazocine (as hydrochloride)

Analgesic for Oral Use—Brief Summary

Indications: For the relief of moderate to severe pain. Contraindication: Talwin should not be administered to patients who are hypersensitive to it.

Warnings: Drug Dependence. There have been instances of psychological and physical dependence on parenteral Talwin in patients with a history of drug abuse and, rarely, in patients without such a history. Abrupt discontinuance following the extended use of parenteral Talwin has resulted in withdrawal symptoms. There have been a few reports of dependence and of withdrawal symptoms with orally administered Talwin. Patients with a history of drug dependence should be under close supervision while receiving Talwin orally.

In prescribing Talwin for chronic use, the physician should take precautions to avoid increases in dose by the patient and to prevent the use of the drug in anticipation of pain rather than for the relief of pain.

Head Injury and Increased Intracranial Pressure. The respiratory depressant effects of Talwin and its potential for elevating cerebrospinal fluid pressure may be markedly exaggerated in the presence of head injury, other intracranial lesions, or a preexisting increase in intracranial pressure. Furthermore, Talwin can produce effects which may obscure the clinical course of patients with head injuries. In such patients, Talwin must be used with extreme caution and only if its use is deemed essential.

Usage in Pregnancy. Safe use of Talwin during pregnancy (other than labor) has not been established. Animal reproduction studies have not demonstrated teratogenic or embryotoxic effects. However, Talwin should be administered to pregnant patients (other than labor) only when, in the judgment of the physician, the potential benefits outweigh the possible hazards. Patients receiving Talwin during labor have experienced no adverse effects other than those that occur with commonly used analgesics. Talwin should be used with caution in women delivering premature infants.

Acute CNS Manifestations. Patients receiving therapeutic doses of Talwin have experienced, in rare instances, hallucinations (usually visual), disorientation, and confusion which have cleared spontaneously within a period of hours. The mechanism of this reaction is not known. Such patients should be very closely observed and vital signs checked. If the drug is reconstituted it should be done with caution since the acute CNS manifestations may recur.

Usage in Children. Because clinical experience in children under 12 years of age is limited, administration of Talwin in this age group is not recommended.

Ambulatory Patients. Since sedation, dizziness, and occasional euphoria have been noted, ambulatory patients should be warned not to operate machinery, drive cars, or unnecessarily expose themselves to hazards.

Precautions: Certain Respiratory Conditions. Although respiratory depression has rarely been reported after oral administration of Talwin, the drug should be administered with caution to patients with respiratory depression from any cause, severe bronchial asthma and other obstructive respiratory conditions, or cyanosis.

Impaired Renal or Hepatic Function. Decreased metabolism of the drug by the liver in extensive liver disease may predispose to accentuation of side effects. Although laboratory tests have not indicated that Talwin causes or increases renal or hepatic impairment, the drug should be administered with caution to patients with such impairment.

Myocardial Infarction. As with all drugs, Talwin should be used with caution in patients with myocardial infarction who have nausea or vomiting.

Biliary Surgery. Until further experience is gained with the effects of Talwin on the sphincter of Oddi, the drug should be used with caution in patients about to undergo surgery of the biliary tract.

Patients Receiving Narcotics. Talwin is a mild narcotic antagonist. Some patients previously given narcotics, including morphine for the daily treatment of narcotic dependence, have experienced mild withdrawal symptoms after receiving Talwin.

CNS Effect. Caution should be used when Talwin is administered to patients prone to seizures; seizures have occurred in a few such patients in association with the use of Talwin although no cause and effect relationship has been established.

Adverse Reactions: Reactions reported after oral administration of Talwin include gastrointestinal: nausea, vomiting; infrequently constipation; and rarely abdominal distress, anorexia, diarrhea. CNS effects: dizziness, lightheadedness, sedation, euphoria, headache; infrequently weakness, disturbed dreams, insomnia, syncope, visual blurring and focusing difficulty, hallucinations (see Acute CNS Manifestations under WARNINGS).

Autonomic: sweating; infrequently tachycardia, hypertension, and rarely chills. Allergic: infrequently rash; and rarely urticaria, edema of the face. Cardiovascular: infrequently decrease in blood pressure, tachycardia. Other: rarely respiratory depression, urinary retention.

Dosage and Administration: Adults. The usual initial adult dose is 1 tablet (50 mg.)

every three or four hours. This may be increased to 2 tablets (100 mg.) when needed.

Total daily dosage should not exceed 600 mg. When antinflammatory or antipyretic effects are desired in addition to analgesia, aspirin can be administered concomitantly with Talwin.

Children Under 12 Years of Age. Since clinical experience in children under 12 years of age is limited, administration of Talwin in this age group is not recommended.

Duration of Therapy. Patients with chronic pain who have received Talwin orally for prolonged periods have not experienced withdrawal symptoms even when administration was abruptly discontinued (see WARNINGS). No tolerance to the analgesic effect has been observed. Laboratory tests of blood and urine and of liver and kidney function have revealed no significant abnormalities after prolonged administration of Talwin.

Overdosage: Manifestations. Clinical experience with Talwin overdosage has been insufficient to define the signs of this condition.

Treatment. Oxygen, intravenous fluids, vasopressors, and other supportive measures should be employed as indicated. Assisted or controlled ventilation should also be considered. Although nalorphine and levorphan are not effective antitoxins for respiratory depression due to overdosage or unusual sensitivity to Talwin, parenteral naloxone (Narcant, available through Endo Laboratories) is a specific and effective antagonist.

Talwin is not subject to narcotic controls.

How Supplied: Tablets, peach color, scored. Each tablet contains Talwin (brand of pentazocine) as hydrochloride equivalent to 50 mg. base. Bottles of 100.

Winthrop Laboratories, New York, N.Y. 10016



50mg. Tablets **Talwin®**  
brand of  
pentazocine  
(as hydrochloride)  
in moderate to severe pain

## Chemotherapy Tested for Joint Disorders



Investigators at the University of California San Diego School of Medicine are assessing the potential of chemical treatment for preventing or reducing joint immobility. Studying data on the tissue changes that surround stiff joints are, left to right, Dr. Wayne Akeson, Professor of Surgery and head of the Division of Orthopedics, and research associates Savio Woo, Ph.D., and David Amiel.

## Researchers Report Progress In Altering Genetic Material

### Medical Tribune Report

WASHINGTON

—

At concurrent sessions of the American Association for the Advancement of Science here, a biochemist from the National Institutes of Health was outlining the difficult problems that lie ahead in altering genetic material for treatment of inborn disease, while a colleague from the University of Maryland was reporting some initial success with a new technique for introducing foreign DNA into cells in tissue culture.

Despite his doubts about the immediate future of "gene therapy"—not to mention his doubts about how society will regard it—Dr. Martin said: "Enormous good will come from further genetic research. Good in areas not necessarily related to inborn errors of metabolism but very possibly in afflictions like cancer and heart disease. I would continue this research at a slow but steady pace."

genes must begin in utero in order to prevent deleterious effects.

• Finally, the DNA or RNA introduced will not only have to be mammalian if it is not to be rejected by the host cells, but will probably have to be human as well.

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## Frozen Marrow Cells May Retain Capacity To Yield Hemoglobin

### Medical Tribune Report

BETHESDA, Md.—Human bone marrow cells stored in the frozen state as long as nine months are able to function normally in the production of hemoglobin according to investigators whose work was supported by the National Institutes of Health.

This finding, the NIH reported, brings closer the day when an individual's own previously frozen and stored marrow cells might be used to reconstitute his production of blood cells following lethal radiation or a catastrophic illness.

The capacity of such stored marrow cells to repopulate the marrow space, it noted, had been demonstrated previously in rodents, dogs, and monkeys.

Dr. John W. Adamson and Rainer Storb, of the University of Washington School of Medicine and Veterans Administration Hospital, Seattle, conducted the new studies. They tested the viability of frozen stored human bone marrow cells by determining their capacity to synthesize hemoglobin in response to treatment with erythropoietin.

In 10 laboratory studies performed on marrow from six individuals, the investigators found that hemoglobin synthesis in treated cultures was increased many times over that of untreated control cultures. Since hemoglobin synthesis takes place only in dividing and growing cells, this observation constitutes evidence that the stored cells do in fact proliferate.

The investigators cautioned that their results apply to only one of the five types of precursor or "stem" cells in the marrow.

The studies received support from the National Cancer Institute, the National Heart and Lung Institute, and the National Institute of Allergy and Infectious Diseases.

# At 10:17a.m. Emmy Burns' future started looking brighter

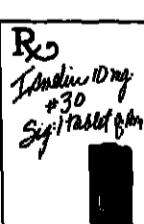


An important step was taken to re-control her hypertension and decrease her vulnerability to organ damage

Emmy Burns just received her prescription for Ismelin. Her blood pressure was no longer responsive to milder agents. So her physician decided that this was the right time to add Ismelin. Because Ismelin is guanethidine, perhaps the most effective anti-hypertensive ever available for moderate to severe hypertension. And when blood pressure is controlled with Ismelin, it usually stays controlled.

## Ismelin® sulfate (guanethidine sulfate)

sooner may  
be better for  
the uncontrolled  
hypertensive



When Ismelin is added to thiazides, increments must be gradual and dosage of all drugs reduced to lowest effective level once blood-pressure control is established.

With reduction of dosage, side effects often are minimized.

Patients should be warned about orthostatic hypotension, especially during initial dosage adjustment and with postural changes. They should avoid sudden or prolonged standing or exercise and should sit or lie down if dizzy or weak.

Uncontrolled hypertension of any degree poses an unacceptable risk to the patient's future well-being.

**ISMELIN® sulfate  
(guanethidine sulfate)**  
**INDICATIONS:** Primarily for severe or sustained elevation of blood pressure (particularly diastolic) in almost all forms of fixed and progressive hypertension, even when blood pressure elevation is moderate. Not recommended for labile or transient hypertension.

**CONTRAINDICATIONS:** Proven or suspected phaeochromocytoma; hypersensitivity to Ismelin. Do not use with MAO inhibitors.

**WARNINGS:** Ismelin is a potent drug and can lead to disturbing and serious clinical problems. Warn patients not to deviate from instructions and about the possible side effects of Ismelin, which can occur frequently. To prevent dizziness, patients should sit or lie down with onset of dizziness or weakness, which may be particularly bothersome during initial dosage adjustment and with postural changes. Prolonged hypotension is most likely in the elderly and is accentuated by hot weather, alcohol, or exercise. Warn patients to avoid sudden or prolonged standing or exercise while taking Ismelin.

Concurrent use with rauwolfia derivatives may cause excessive postural hypotension, bradycardia, and/or syncope. If possible, withdraw therapy 2 weeks prior to surgery to avoid possible vascular collapse and to reduce hazard of cardiac arrest during anesthesia. If emergency surgery is indicated, administer preanesthetic and anesthetic agents cautiously in reduced doses with oxygen, atropine, and vasopressors ready for immediate use. Patients with arrhythmias during initial dosage of Ismelin may have a greater propensity for cardiac arrhythmias.

Refractory hypertension may reduce dosage requirements. In frank congestive heart failure due to hypertension, Ismelin is not recommended. Do not use in patients with (a) hypotension; (b) catecholamine depletion and increased responsiveness to norepinephrine; special care is required when treating patients with a history of bronchial asthma, since this condition may be aggravated.

**Use in Pregnancy:**

The safety of Ismelin for use in pregnancy has not been established; therefore, this drug should be used in pregnant patients only when, in the judgment of the physician, its use is deemed essential to the welfare of the patient.

**PRECAUTIONS:** Give very cautiously to hypertension with (a) renal disease with nitrogen retention; (b) congestive heart failure; (c) peripheral vascular disease, especially with encopresis; and (d) rising BUN levels. Give with extreme caution to those with severe congestive failure. Watch for weight gain or edema. If possible, withdraw therapy. If Ismelin is used with digitalis, remember that both drugs slow the heart rate.

Antihypertensive drugs (eg, angiotensinase, mild stimulants (eg, epinephrine, methylphenidate), and tricyclic antidepressants (eg, imipramine, propantheline) may increase hypotension.

Use of Ismelin may cause orthostatic hypotension or syncope.

When such measures fail, psychoanalytic therapy may be helpful in finding the cause of

hypotension. The woman sleeps in a separate bed or separate bedroom. The family has one room, two rooms. The children are sleeping in the same room. Grandparents are living in the home. You don't get all these cues in an interview. Either they forget to tell you or they are embarrassed.

In treating insomnia, Dr. Karacan believes that drugs should be used only as a last resort. In fact, he remarked, some patients are already "walking pharmacists"; administer eight or 10 drugs to

themselves daily, "and if you simply take

away all the drugs they are already taking, the insomnia leaves with the drugs."

"There's no question that at least 60 to

70 per cent of the self-defined insomniacs

could be cured of their problem without

drugs," he said. "But sometimes it takes a

bit of time to find out what the real prob-

lem is. You don't often find it in a five-

minute consultation, and general practi-

tioners have very little time to talk over

the problems of such patients."

For many insomniacs, Dr. Karacan con-

tinued, some changes in life style or

eating and drinking habits prove to be a

cure. For example, if a low arousal thresh-

old or something else in the arousal sys-

tem seems to be the cause of the insomnia,

he recommends a quieter life, with avoid-

ance of alcohol and parties and no watch-

ing of TV or reading or exciting novels

before bedtime.

When, as a last resort Dr. Karacan said,

gives a drug for the sleeping problem itself,

he gives it in a pattern of five nights on

the drug and two nights off it.

"So the patient doesn't sleep for two

nights," he commented. "It's better than

not sleeping every night and better than

becoming addicted to hypnotics."

He concluded: "Insomnia is a hetero-

geneous group, there isn't one type of

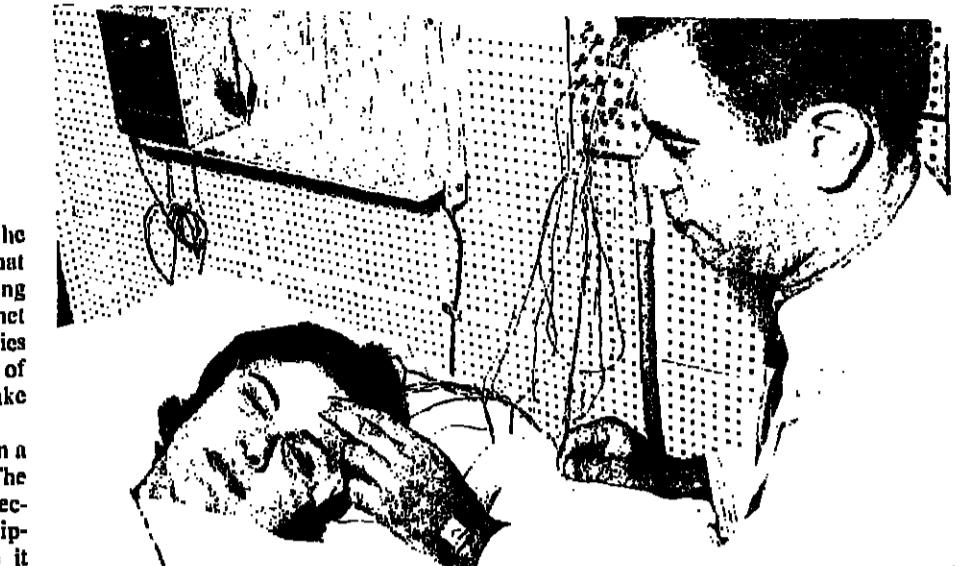
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sultation and a prescription for a sleeping

pill just doesn't work."

In this exclusive roundup MEDICAL TRIBUNE is publishing highlights from the First European Congress on Sleep Research, held in Basel, Switzerland.



## Insomnia Study Is Facilitated By Mobile Unit

Medical Tribune World Service

BASEL, SWITZERLAND—Convinced that he could often learn much more quickly what was causing a case of insomnia by making studies in the patient's home, Dr. Ismet Karacan, director of Sleep Laboratories at the University of Florida College of Medicine, has set up a mobile unit to take the laboratory to the patient.

The equipment truck is parked within a mile radius of the patient's home. The doctor visits the patient, puts the electrodes on his head, gives him an equipment activator, and tells him to use it when he wants to go to sleep.

"The patient laboratory can contaminate the data," Dr. Karacan told MEDICAL TRIBUNE. "You bring the subject into another social environment, an artificial environment. . . . I want to see the patient in his own environment."

away all the drugs they are already taking, the insomnia leaves with the drugs."

"There's no question that at least 60 to 70 per cent of the self-defined insomniacs could be cured of their problem without drugs," he said. "But sometimes it takes a bit of time to find out what the real problem is. You don't often find it in a five-minute consultation, and general practitioners have very little time to talk over the problems of such patients."

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## Hypersomnia: Third Variety Said to Exist

Medical Tribune World Service

Hypersomnias have generally been classified into two types—those characterized by non-REM sleep periods and those in which the patient has both non-REM and REM sleep in a normal pattern but repeats the pattern over a longer period than normal.

A case study indicating that still another type of hypersomnia exists was presented to the Sleep Congress by Drs. R. Broughton and A. Guzman, of the University of Ottawa's Departments of Medicine and Pharmacology.

DR. GUZMAN

The third type, they said, is a REM hypersomnia, and it is improved by REM suppressives. Imipramine cured their patient, an 18-year-old boy, apparently permanently, they reported.

## Temperature for Sleeping Is Best From 27 to 36° C.

Medical Tribune World Service

The optimal range of temperature for restful sleep is between 27° and 36° C., and the most comfortable temperature for sleeping is at the lower end of this range, according to two sleep investigators at the Neurologische Universitätsklinik mit Abteilung für Neurophysiologie, Freiburg, West Germany.

Drs. K. Kendel and W. Schmidt-Kessen said all results obtained thus far on the climatic influence on sleep had been related to extreme experimental conditions. No one had tested the influence of conditions as near normal as possible on the restful sleep of normal young adults.

Undressed Subjects Shivered

Poligraphically recording the night sleep of normal male students at varying room temperatures, they found that undressed and uncovered subjects began shivering from cold just 1° below the temperature for most comfortable rest, 27°. More than 10° higher, above 37°, they began profuse sweating and reported having unpleasant heat rashes.

Among the other findings, they noted that the higher the room temperature, the more restless the sleeper, and that the heart beat went up with room temperature. On cooler nights the subjects had more REM sleep, but also, their remembrance of dreams was lower.

of medicines was the direct result of hospitalization. Why don't patients take medicines properly? The study cited such reasons as: they forget, they can't keep track of all of them they are supposed to take, they feel better (or worse), they used them up, friends said they were dangerous (or worthless). What can we do about this problem? If possible, reduce polypharmacy, switch medicines to be taken only once daily, provide better information to the patient and his relatives (preferably written), and try to make the patient bring along what remains of his medicine at his next doctor's visit. Industry can help by using throwaway packaging of various types, or packaging with calendars, as for contraceptive pills. Perhaps half our patients take medicines in ways other than those prescribed. It's up to each one of us to find the solution for our own patients. C. F. Borchgrevink, editorial, *Tidsskrift for den Norske Lægeforening* [J. Norwegian M. A.] 92:34, December 10, 1972.

**Editorial Capsule**  
... brief summaries of editorials or guest editorials in current medical journals.

**Daze of Retirement**  
Physicians should be "deeply concerned with policies that call for arbitrary retirement based on chronologic age, without regard to individual desires or capabilities."

**Progress in Leprosy**  
About 25 years ago, most studies on leprosy were performed by "dedicated workers, as isolated as their patients; communication was a formidable task and fraught with language difficulties. . . . However in the latter part of the 1950's and the early 1960's scientists, as distinct from humanitarian-oriented field workers, began to take an interest in the problems of leprosy. Microbiologists, statisticians, immunologists, epidemiologists and re-

C I B A